FILE CAPLUS

Claim 1

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=> d .ca 18 21-50

ANSWER 21 OF 86 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2000:875755 CAPLUS

DOCUMENT NUMBER:

134:41092

TITLE:

Compounds and methods for treatment and diagnosis of

mycobacterial infections

INVENTOR(S):

Visser, Elizabeth

PATENT ASSIGNEE(S):

Genesis Research and Development Corporation Limited,

N.Z.

SOURCE:

U.S., 147 pp., Cont.-in-part of U.S. 5,985,287.

CODEN: USXXAM

DOCUMENT TYPE:

Patent English

LANGUAGE: FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND DATE	APPLICATION NO.	DATE
00 010000	A 20001212	US 1998-95855	
US 6284255			
	A 19991214	00 200 0 0 0 0 0 0	
US 5985287			
US 6406704		US 1998-205426	
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	A2 19990701	WO 1998-NZ189	19981223 <
	A3 19991202		
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		AU 1999-18936	19981223 <
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BR 9814432		BR 1998-14432	19981223 <
		EP 1998-963665	

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                                            IN 2000-CA231
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    NO 2000003261
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                                            NO 2000-3261
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PRIORITY APPLN. INFO.:
                                            US 1996-705347
                                                                A2 19960829
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                                                                A2 19970612
                                            US 1997-997362
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                                            US 1997-996624
                                                                A 19971223
                                            US 1997-997080
                                                                A 19971223
                                                                A 19980216
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                                            US 1998-95855
                                                                A2 19980611
                                            US 1998-156181
                                                                A 19980917
                                            US 1998-205426
                                                                A 19981204
                                            WO 1998-NZ189
                                                                W 19981223
ED
    Entered STN: 14 Dec 2000
    The present invention provides polypeptides comprising an immunogenic
ΔR
    portion of a M. vaccae protein and DNA mols. encoding such polypeptides,
     together with methods for their use in the diagnosis and treatment of
    mycobacterial infection. Methods for enhancing the immune response to an
    antigen including administration of M. vaccae culture filtrate,
    delipidated M. vaccae cells or delipidated and deglycolipidated M. vaccae
     cells are also provided.
IC
     ICM C07K014-35
     ICS A61K039-04
INCL 530350000
CC
     15-2 (Immunochemistry)
     Section cross-reference(s): 3, 9
IT
    Blood analysis
    Blood plasma
    Blood serum
    DNA sequences
     Immunotherapy
     Infection
    Macrophage
    Molecular cloning
    Mycobacterium
    Mycobacterium avium
    Mycobacterium tuberculosis
    Mycobacterium vaccae
    Neoplasm
      Protein sequences
     Skin
    Tuberculosis
    Urine analysis
    Vaccines
        (Mycobacterium vaccae antigens as vaccine, diagnostic,
        immunotherapeutic and adjuvant for infection, immunol. disease, and
        cancer)
    Fusion proteins (chimeric proteins)
IT
    RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (Mycobacterium vaccae antigens as vaccine, diagnostic,
        immunotherapeutic and adjuvant for infection, immunol. disease, and
        cancer)
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                   204785-82-0 204785-84-2, Antigen GV-24 (Mycobacterium
     204785-67-1
               204785-86-4, Antigen GV-25 (Mycobacterium vaccae)
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Antigen GV-27 (Mycobacterium vaccae N-terminal fragment)
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    Antigen GV-27B (Mycobacterium vaccae fragment)
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    GV-22B (Mycobacterium vaccae)
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                                           228107-96-8, Antigen GVs-9
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    (Mycobacterium vaccae)
    228108-01-8, Antigen GV-45 (Mycobacterium vaccae) 228108-08-5, Antigen
                                    228108-12-1, Antigen GV-44 (Mycobacterium
    GV-41B (Mycobacterium vaccae)
              228108-14-3 250277-71-5
    vaccae)
    RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
     (Biological study)
        (amino acid sequence; Mycobacterium vaccae antigens as vaccine,
       diagnostic, immunotherapeutic and adjuvant for infection, immunol.
       disease, and cancer)
    119939-22-9, Antigen α (Mycobacterium BCG clone pαL-1
                         136108-01-5, Antigen (Mycobacterium tuberculosis
    precursor reduced)
    32.0-kilodalton precursor reduced)
                                        145110-35-6, Antigen 85C
     (Mycobacterium tuberculosis clone 11-2 precursor)
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    85B (Mycobacterium leprae precursor reduced)
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    204785-69-3, Antigen GVc-7 (Mycobacterium vaccae)
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    204786-07-2
                  204786-10-7 204786-11-8 204786-15-2
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    228105-87-1, Antigen GVc-13 (Mycobacterium vaccae) 228107-33-3, Antigen
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    GV-1/70 (Mycobacterium vaccae)
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    RL: PRP (Properties)
        (unclaimed protein sequence; compds. and methods
       for treatment and diagnosis of mycobacterial infections)
                              THERE ARE 19 CITED REFERENCES AVAILABLE FOR THIS
REFERENCE COUNT:
                        19
                              RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
    ANSWER 22 OF 86 CAPLUS COPYRIGHT 2006 ACS on STN
                        2000:861814 CAPLUS
ACCESSION NUMBER:
                        134:26782
DOCUMENT NUMBER:
                        Identification and characterization of Snf2 related
TITLE:
                        CBP activator protein (SRCAP)
                        Chrivia, John; Yaciuk, Peter
INVENTOR(S):
                        Saint Louis University, USA
PATENT ASSIGNEE(S):
                        PCT Int. Appl., 103 pp.
SOURCE:
                        CODEN: PIXXD2
DOCUMENT TYPE:
                        Patent
LANGUAGE:
                        English
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
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IT

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PATENT NO.
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                                         APPLICATION NO.
                                                                DATE
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PRIORITY APPLN. INFO.:
                                           US 1999-136620P
                                                               P 19990527
                                           US 2000-579181
                                                               A 20000525
    Entered STN: 08 Dec 2000
ED
AB
    A protein, SRCAP, a novel SNF2/SWI2 protein family member interacting with
    CREB-binding proteins, is provided. The protein is capable of
    co-activating CREB binding protein (CBP) mediated transcription, as well
    as activating transcription without CBP. SRCAP is a Snf2 family member.
    As such, it has ATPase activity. Consistent with this activity, SRCAP
    contains the conserved ATPase domain found within members of the Snf2
    family. Transfection expts. demonstrate that SRCAP is able to activate
    transcription when expressed as a Gal-SRCAP chimera and that SRCAP also
    enhances the ability of CBP to activate transcription. The adenoviral
    protein E1A is found to disrupt interaction between SRCAP and CBP possibly
    representing a mechanism for E1A-mediated transcriptional repression.
    SRCAP also interacts with NS5A of hepatitis C virus (HCV), and this
    interaction may have effect on growth regulation of cells infected with
    HCV through the down-regulation of p21 promoter activity and contribute
    HCV pathogenesis. Fragments of SRCAP are also provided, as are its cDNA
    and cDNA fragments. Antibodies that bind to SRCAP are also provided.
    ICM C12N015-52
TC
         C12N015-62; C12N015-11; C12N005-10; C12N009-14; C07K016-40;
    ICS
         G01N033-50; G01N033-566; C12Q001-68; A61K038-46
CC
    6-3 (General Biochemistry)
    Section cross-reference(s): 3, 13
IT
    Transcription factors
    RL: BPR (Biological process); BSU (Biological study, unclassified); BUU
     (Biological use, unclassified); BIOL (Biological study); PROC (Process);
    USES (Uses)
        (GAL4, fusion product with SRCAP; cloning and sequence of
       transcription factor SRCAP which interacts with CREB-binding protein)
IT
    Protein sequences
    cDNA sequences
        (identification and characterization of Snf2 related CBP activator
       protein (SRCAP))
IT
    Fusion proteins (chimeric proteins)
    RL: BPR (Biological process); BSU (Biological study, unclassified); BUU
     (Biological use, unclassified); BIOL (Biological study); PROC (Process);
    USES (Uses)
        (of GAL4 and SRCAP; cloning and sequence of transcription factor SRCAP
       which interacts with CREB-binding protein)
    311824-07-4 311824-09-6
TT
    RL: BPR (Biological process); BSU (Biological study, unclassified); PRP
     (Properties); BIOL (Biological study); PROC (Process)
        (amino acid sequence; cloning and sequence of transcription factor
       SRCAP which interacts with CREB-binding protein)
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THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS

REFERENCE COUNT:

RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

ANSWER 23 OF 86 CAPLUS COPYRIGHT 2006 ACS on STN L8

2000:861803 CAPLUS ACCESSION NUMBER:

DOCUMENT NUMBER: 134:26110

TITLE: Human secreted and transmembrane polypeptides and

nucleic acids encoding the same

Ashkenazi, Avi J.; Baker, Kevin P.; Botstein, David; INVENTOR(S):

Desnoyers, Luc; Eaton, Dan L.; Ferrara, Napoleone; Fong, Sherman; Gerber, Hanspeter; Gerritsen, Mary E.;

Goddard, Audrey; Godowski, Paul J.; Grimaldi, Christopher J.; Gurney, Austin L.; Kljavin, Ivar J.; Napier, Mary A.; Pan, James; Paoni, Nicholas F.; Roy, Margaret Ann; Stewart, Timothy A.; Tumas, Daniel;

Watanabe, Colin K.; Williams, P. Mickey; Wood, William

I.; Zhang, Zemin Genentech, Inc., USA PATENT ASSIGNEE(S): PCT Int. Appl., 935 pp. SOURCE:

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English FAMILY ACC. NUM. COUNT: 143

PATENT INFORMATION:

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WO	2000				A1		2000	1207		WO 2000-US8439										
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WO	9963	880			A2		1999	1209		WO 1	999-1	US12	252		1	9990	602 <-			
WO	9963	088			A3		2001	0329												
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ΕP	1210	418			A1		2002	0605		EP 2	000-	9166	75		2	0000	330			
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AU 2000069018
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ED Entered STN: 08 Dec 2000

The present invention is directed to novel polypeptides and to nucleic acid mols. encoding those polypeptides. Thus, 135 cDNA sequences encoding human secreted and/or transmembrane proteins are identified by extracellular domain homol. screening, amylase screening, and a signal sequence algorithm to identify novel polypeptides. The proteins exhibit various biol. activities useful for diagnostic and therapeutic applications. Also provided herein are vectors and host cells comprising those nucleic acid sequences, chimeric polypeptide mols. comprising the polypeptides of the present invention fused to heterologous polypeptide sequences, antibodies which bind to the polypeptides of the present invention and to methods for producing the polypeptides of the present invention.

IC ICM C12N015-12 ICS C07K014-47; C07K014-705; C12N015-62; C07K016-18; G01N033-53;

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                               THERE ARE 11 CITED REFERENCES AVAILABLE FOR THIS
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ANSWER 24 OF 86 CAPLUS COPYRIGHT 2006 ACS on STN
                        2000:772766 CAPLUS
ACCESSION NUMBER:
DOCUMENT NUMBER:
                        133:330556
                        Genome sequence and polypeptides of Pyrococcus abyssi
TITLE:
                        and their uses
                        Forterre, Patrick; Thierry, Jean-Claude; Prieur,
INVENTOR(S):
                        Daniel; Dietrich, Jacques; Lecompte, Odile; Querellou,
                        Joel; Weissenbach, Jean; Saurin, William; Heilig,
                        Roland; Flament, Didier; Raffin, Jean-Paul; Henneke,
                        Ghislaine; Gueguen, Yannick; Rolland, Jean-Luc
PATENT ASSIGNEE(S):
                        Centre National de la Recherche Scientifique (CNRS),
                        Fr.; Institut Français de Recherche pour
                        l'Exploitation de la Mer - IFREMER
                        PCT Int. Appl., 1403 pp.
SOURCE:
                        CODEN: PIXXD2
DOCUMENT TYPE:
                        Patent
LANGUAGE:
                        French
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
     PATENT NO.
                      KIND DATE
                                         APPLICATION NO.
                                                               DATE
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            LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE,
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PRIORITY APPLN. INFO.:
                                           FR 1999-5034
                                                              A 19990421
                                           WO 2000-FR1065
                                                              W 20000421
    Entered STN: 03 Nov 2000
ED
    The invention relates to the genome sequence of Pyrococcus abyssi strain
AΒ
    Orsay, the 807 open reading frame nucleotide sequences coding for
    polypeptides of P. abyssi such as polypeptides involved in metabolism or in
    the replication process, in addition to vectors including said sequences and
    cells transformed by said vectors. Replication factor C (large and small
    forms resulting from intein splicing), PCNA (proliferating cell nuclear
    antigen), DNA polymerase II large and small subunits, replication factor
    A, and DNA polymerase I were isolated and characterized by recombinant
    cloning in Escherichia coli. The invention also relates to methods using
     said nucleic acids or polypeptides, especially biosynthesis methods or
    biodegrdn. methods for mols. of interest and to kits comprising said
    polypeptides.
IC
    ICM C12N015-31
         C12N015-54; C12N015-55; C12N015-57; C12N015-60; C12N015-61;
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     Genome
     Molecular cloning
       Protein sequences
     Pyrococcus abyssi
        (genome sequence and polypeptides of Pyrococcus abyssi and their uses)
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     RL: BPN (Biosynthetic preparation); BUU (Biological use, unclassified);
     BIOL (Biological study); PREP (Preparation); USES (Uses)
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   (amino acid sequence; genome sequence and polypeptides of Pyrococcus
   abyssi and their uses)
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L8 ANSWER 25 OF 86 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2000:742133 CAPLUS

DOCUMENT NUMBER: 133:291993

TITLE: Cloning and cDNA and deduced amino acid sequences of

48 human secreted proteins

INVENTOR(S): Rosen, Craig A.; Ruben, Steven M.; Komatsoulis, George

PATENT ASSIGNEE(S): Human Genome Sciences, Inc., USA

SOURCE: PCT Int. Appl., 500 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 4

PATENT INFORMATION:

PAT	PATENT NO.									APPLICATION NO.							
WO	2000	 0616	 25		A1 20001019			WO 2000-US8981									
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Entered STN: 20 Oct 2000
ED
     The present invention relates to 48 novel human secreted proteins and
AB
     isolated nucleic acids containing the coding regions of the genes encoding
     such proteins. Tissue distribution, sequence homologies, and preferred
     epitope sites are provided for the secreted proteins, as well as
     chromosomal mapping of some of the genes. Also provided are vectors, host
     cells, antibodies, and recombinant methods for producing human secreted
     proteins in bacterial, insect, and mammalian cells. The invention further
     relates to diagnostic and therapeutic methods useful for diagnosing and
     treating disorders related to these novel human secreted proteins.
     High-throughput screening assays are also provided for various putative
     activities of the secreted proteins.
     ICM C07K014-47
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     3-3 (Biochemical Genetics)
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     Section cross-reference(s): 6, 13, 63
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        (fusion products; cloning and cDNA and deduced amino acid
        sequences of 48 human secreted proteins)
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ACCESSION NUMBER:
                         2000:742132 CAPLUS
DOCUMENT NUMBER:
                         133:291992
                         Cloning and cDNA and deduced amino acid sequences of
TITLE:
                         48 human secreted proteins
                         Rosen, Craig A.; Ruben, Steven M.; Komatsoulis, George
INVENTOR(S):
PATENT ASSIGNEE(S):
                         Human Genome Sciences, Inc., USA
SOURCE:
                         PCT Int. Appl., 478 pp.
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CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

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ED Entered STN: 20 Oct 2000

AB The present invention relates to 48 novel human secreted proteins and isolated nucleic acids containing the coding regions of the genes encoding such proteins. Tissue distribution, sequence homologies, and preferred epitope sites are provided for the secreted proteins, as well as chromosomal mapping of some of the genes. Also provided are vectors, host cells, antibodies, and recombinant methods for producing human secreted proteins in bacterial, insect, and mammalian cells. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and treating disorders related to these novel human secreted proteins. High-throughput screening assays are also provided for various putative activities of the secreted proteins.

IC ICM C07K014-47

ICS C07K015-52; C12N005-10; C12N005-16; C12N015-12; C12N015-19; C12N015-63; C12N015-64

CC 3-3 (Biochemical Genetics)

Section cross-reference(s): 6, 13, 63

IT Immunoglobulins

RL: BPN (Biosynthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(fusion products; cloning and cDNA and deduced amino acid sequences of 48 human secreted proteins)

IT Protein sequences

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(of 48 human secreted proteins)
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RL: BOC (Biological occurrence); BPN (Biosynthetic preparation); BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); OCCU (Occurrence); PREP (Preparation); USES (Uses)

(amino acid sequence; cloning and cDNA and deduced amino acid sequences of 48 human secreted proteins)

TΤ 300625-48-3 300625-49-4 300625-50-7 300625-51-8 300625-52-9 300625-56-3 300625-57-4 300625-53-0 300625-54-1 300625-55-2 300625-58-5 300625-59-6 300625-60-9 300625-61-0 300625-62-1 300625-63-2 300625-64-3 300625-65-4 300625-66-5 300625-67-6 300625-68-7 300625-69-8 300625-70-1 300625-71-2 300625-72-3 300625-73-4 300681-86-1 300697-20-5 300697-36-3 300697-49-8 300697-70-5 300701-81-9 300702-03-8

RL: PRP (Properties)

(unclaimed protein sequence; cloning and cDNA and

deduced amino acid sequences of 48 human secreted proteins)

REFERENCE COUNT: 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 27 OF 86 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2000:718218 CAPLUS

DOCUMENT NUMBER: 133:291969

TITLE: Viral encoded semaphorin protein receptor (VESPR) cDNA

and its polypeptides

INVENTOR(S): Spriggs, Melanie K.; Comeau, Michael R.; Dubose,

Robert F.; Johnson, Richard S.

PATENT ASSIGNEE(S): Immunex Corporation, USA

SOURCE: U.S., 32 pp.
CODEN: USXXAM

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE			
US 6130068	Α	20001010	US 1998-181706		19981028 <		
US 6174689	B1	20010116	US 1999-458791		19991210		
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US 6562949	B1	20030513	US 1999-459065		19991210		
US 2003095968	A1	20030522	US 2002-294055		20021113		
PRIORITY APPLN. INFO.:			US 1997-112009P	P	19971028		
			US 1997-958598	Α	19971028		
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ED Entered STN: 11 Oct 2000

AB The invention is directed to VESPR polypeptides as a purified and isolated protein, the DNA encoding the VESPR polypeptide, host cells transfected with cDNAs encoding VESPR, and methods for preparing VESPR polypeptides.

IC ICM C07H021-04

ICS C12N015-09; C12N015-63

INCL 435069100

CC 3-3 (Biochemical Genetics)

Section cross-reference(s): 1, 6, 14

IT Fusion proteins (chimeric proteins)

RL: BPN (Biosynthetic preparation); BIOL (Biological study); PREP (Preparation)

(of VESPR; viral encoded semaphorin protein receptor (VESPR) cDNA and polypeptides)

IT Protein sequences

(of human VESPR; viral encoded semaphorin protein receptor (VESPR) cDNA and polypeptides) 209215-07-6DP, subfragments claimed 223775-30-2P IT 223775-32-4P RL: BPN (Biosynthetic preparation); BPR (Biological process); BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); PROC (Process); USES (Uses) (amino acid sequence; properties and therapeutic uses of VESPR (Viral Encoded Semaphorin Protein Receptor)) 224041-58-1, Semaphorin A39R (Ectromelia virus) IT RL: PRP (Properties) (unclaimed protein sequence; viral encoded semaphorin protein receptor (VESPR) cDNA and its polypeptides) THERE ARE 23 CITED REFERENCES AVAILABLE FOR THIS REFERENCE COUNT: 23 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT ANSWER 28 OF 86 CAPLUS COPYRIGHT 2006 ACS on STN 2000:707187 CAPLUS ACCESSION NUMBER: DOCUMENT NUMBER: 133:262326 Cloning and cDNA and deduced amino acid sequences of TITLE: 47 human secreted proteins Rosen, Craig A.; Ruben, Steven M.; Komatsoulis, George INVENTOR (S): Human Genome Sciences, Inc., USA PATENT ASSIGNEE(S): PCT Int. Appl., 387 pp. SOURCE: CODEN: PIXXD2 DOCUMENT TYPE: Patent English LANGUAGE: FAMILY ACC. NUM. COUNT: PATENT INFORMATION: APPLICATION NO. KIND DATE DATE PATENT NO. -----A1 20001005 WO 2000-US7534 _____ ----WO 2000058335 20000322 <--W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM,

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ED Entered STN: 06 Oct 2000

AB The present invention relates to 47 novel human secreted proteins and isolated nucleic acids containing the coding regions of the genes encoding such proteins. Tissue distribution, sequence homologies, and preferred epitope sites are provided for the secreted proteins, as well as chromosomal mapping of some of the genes. Also provided are vectors, host cells, antibodies, and recombinant methods for producing human secreted proteins in bacterial, insect, and mammalian cells. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and treating disorders related to these novel human secreted proteins.

High-throughput screening assays are also provided for various putative activities of the secreted proteins. IC ICM C07H021-04 ICS C07K014-00; C07K016-00; C12N015-00; C12N015-63; C12N015-85; C12N015-86; C12Q001-68; G01N033-53 3-3 (Biochemical Genetics) CC Section cross-reference(s): 6, 13, 63 Immunoglobulins IT RL: BPN (Biosynthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses) (fusion products; cloning and cDNA and deduced amino acid sequences of 47 human secreted proteins) IT Protein sequences (of 47 human secreted proteins) 295827-95-1 295827-96-2 IT 195890-61-0 295827-97-3 295827-98-4 **295827-99-5** 295828-00-1 295828-01-2 295828-02-3 295828-05-6 295828-06-7 295828-03-4 295828-04-5 295828-07-8 295828-08-9 295828-09-0 295828-74-9 296251-99-5 296267-99-7 296268-00-3 296268-01-4 296268-02-5 296268-03-6 296268-04-7 296268-05-8 296268-06-9 296268-07-0 296268-08-1 296268-10-5 RL: PRP (Properties) (unclaimed protein sequence; cloning and cDNA and deduced amino acid sequences of 47 human secreted proteins) THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS REFERENCE COUNT: 4 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT ANSWER 29 OF 86 CAPLUS COPYRIGHT 2006 ACS on STN ACCESSION NUMBER: 2000:688363 CAPLUS DOCUMENT NUMBER: 133:248090 Cloning and cDNA and deduced amino acid sequences of TITLE: 48 human secreted proteins INVENTOR(S): Rosen, Craig A.; Ruben, Steven M.; Komatsoulis, George PATENT ASSIGNEE(S): Human Genome Sciences, Inc., USA SOURCE: PCT Int. Appl., 407 pp. CODEN: PIXXD2 DOCUMENT TYPE: Patent LANGUAGE: English FAMILY ACC. NUM. COUNT: PATENT INFORMATION: PATENT NO. KIND DATE APPLICATION NO. DATE ---------20000928 WO 2000-US6782 WO 2000056881 A1 W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE,

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     Entered STN: 29 Sep 2000
     The present invention relates to 48 novel human secreted proteins and
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     isolated nucleic acids containing the coding regions of the genes encoding
     such proteins. Tissue distribution, sequence homologies, and preferred
     epitope sites are provided for the secreted proteins, as well as
     chromosomal mapping of some of the genes. Also provided are vectors, host
     cells, antibodies, and recombinant methods for producing human secreted
     proteins in bacterial, insect, and mammalian cells. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and
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     High-throughput screening assays are also provided for various putative
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     Section cross-reference(s): 6, 13, 63
     Immunoqlobulins
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     RL: BPN (Biosynthetic preparation); THU (Therapeutic use); BIOL
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REFERENCE COUNT:
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                                RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
     ANSWER 30 OF 86 CAPLUS COPYRIGHT 2006 ACS on STN
                         2000:688253 CAPLUS
ACCESSION NUMBER:
DOCUMENT NUMBER:
                         133:248082
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TITLE:
                         Cloning and cDNA and deduced amino acid sequences of
                         49 human secreted proteins
INVENTOR(S):
                        Rosen, Craig A.; Ruben, Steven M.; Komatsoulis, George
PATENT ASSIGNEE(S):
                        Human Genome Sciences, Inc., USA
SOURCE:
                         PCT Int. Appl., 431 pp.
                        CODEN: PIXXD2
DOCUMENT TYPE:
                        Patent
LANGUAGE:
                        English
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
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    Entered STN: 29 Sep 2000
ED
    The present invention relates to 49 novel human secreted proteins and
AB
    isolated nucleic acids containing the coding regions of the genes encoding
    such proteins. Tissue distribution, sequence homologies, and preferred
    epitope sites are provided for the secreted proteins, as well as
    chromosomal mapping of some of the genes. Also provided are vectors, host
    cells, antibodies, and recombinant methods for producing human secreted
    proteins in bacterial, insect, and mammalian cells. The invention further
    relates to diagnostic and therapeutic methods useful for diagnosing and
    treating disorders related to these novel human secreted proteins.
    High-throughput screening assays are also provided for various putative
    activities of the secreted proteins.
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    ICM C07H021-04
         C07K014-00; C07K016-00; C12N015-00; C12N015-63; C12N015-85;
    ICS
         C12N015-86; C12Q001-68; G01N033-53
CC
    3-3 (Biochemical Genetics)
    Section cross-reference(s): 6, 13, 63
IT
    Immunoglobulins
    RL: BPN (Biosynthetic preparation); THU (Therapeutic use); BIOL
     (Biological study); PREP (Preparation); USES (Uses)
        (fusion products; cloning and cDNA and deduced amino acid
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    BIOL (Biological study); OCCU (Occurrence); PREP (Preparation); USES
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REFERENCE COUNT:
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ACCESSION NUMBER:
                        2000:666922 CAPLUS
DOCUMENT NUMBER:
                        133:248079
                        Cloning and cDNA and deduced amino acid sequences of
TITLE:
                        27 human secreted proteins
                        Ruben, Steven M.; Ni, Jian; Ebner, Reinhard; Rosen,
INVENTOR(S):
                        Craig A.; Shi, Yanggu; Birse, Charles; Florence,
                        Kimberly; Komatsoulis, George; Lafleur, David W.;
                        Moore, Paul A.; Olsen, Henrik S.; Young, Paul E.
PATENT ASSIGNEE(S):
                        Human Genome Sciences, Inc., USA
                        PCT Int. Appl., 453 pp.
SOURCE:
                        CODEN: PIXXD2
DOCUMENT TYPE:
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LANGUAGE:
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
                              DATE
    PATENT NO.
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ED
     Entered STN: 22 Sep 2000
AB
     The present invention relates to 27 novel human secreted proteins and
     isolated nucleic acids containing the coding regions of the genes encoding
     such proteins. Tissue distribution, sequence homologies, and preferred
     epitope sites are provided for the secreted proteins, as well as
     chromosomal mapping of some of the genes. Also provided are vectors, host
     cells, antibodies, and recombinant methods for producing human secreted
     proteins in bacterial, insect, and mammalian cells. The invention further
     relates to diagnostic and therapeutic methods useful for diagnosing and
     treating disorders related to these novel human secreted proteins.
     High-throughput screening assays are also provided for various putative
     activities of the secreted proteins.
     ICM C12Q001-68
IC
         C07H021-00; C07K005-00
     3-3 (Biochemical Genetics)
CC
     Section cross-reference(s): 6, 13, 63
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     Immunoglobulins
     RL: BPN (Biosynthetic preparation); THU (Therapeutic use); BIOL
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REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 32 OF 86 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2000:666771 CAPLUS

DOCUMENT NUMBER: 133:233613

TITLE: Cloning and cDNA and deduced amino acid sequences of

50 human secreted proteins

INVENTOR(S): Rosen, Craig A.; Ruben, Steven M.; Komatsoulis, George

PATENT ASSIGNEE(S): Human Genome Sciences, Inc., USA

SOURCE: PCT Int. Appl., 453 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

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ED Entered STN: 22 Sep 2000

- The present invention relates to 50 novel human secreted proteins and isolated nucleic acids containing the coding regions of the genes encoding such proteins. Tissue distribution, sequence homologies, and preferred epitope sites are provided for the secreted proteins, as well as chromosomal mapping of some of the genes. Also provided are vectors, host cells, antibodies, and recombinant methods for producing human secreted proteins in bacterial, insect, and mammalian cells. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and treating disorders related to these novel human secreted proteins. High-throughput screening assays are also provided for various putative activities of the secreted proteins.
- IC ICM C07K014-47

ICS C12N005-10; C12N005-16; C12N015-12; C12N015-63; C12N015-64

CC 3-3 (Biochemical Genetics)

Section cross-reference(s): 6, 13, 63

IT Immunoglobulins

RL: BPN (Biosynthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(fusion products; cloning and cDNA and deduced amino acid sequences of 50 human secreted proteins)

IT Protein sequences

(of 50 human secreted proteins)

IT 292103-74-3 292103-75-4 292103-76-5 292103-77-6 292103-78-7

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(unclaimed protein sequence; cloning and cDNA and

deduced amino acid sequences of 50 human secreted proteins)

REFERENCE COUNT: 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 33 OF 86 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2000:646123 CAPLUS

DOCUMENT NUMBER: 133:234221

TITLE: Hermansky pudlak syndrome protein-interacting proteins

(HPSIPs), their complexes with HPS proteins, and

diagnosis and treatment of diseases Nandabalan, Krishnan; Yang, Meijia

INVENTOR(S): Nandabalan, Krishnan; Yang
PATENT ASSIGNEE(S): Curagen Corporation, USA

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SOURCE: PCT Int. Appl., 79 pp.
CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

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	WO 2000053733					A2 20000914			WO 2000-US6518						20000310 <					
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			SI,	SK,	SL,	ТJ,	TM,	TR,	TT,	TZ,	UA,	ŪĠ,	US,	UΖ,	VN,	ΥU,	ZA,	ZW,		
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ED Entered STN: 15 Sep 2000

AB Provided are complexes of Hermansky pudlak syndrome (HPS) proteins and HPS-interacting proteins (HPSIPs). The HPSIPs include 14-3-3 protein, Hrs, atrophin-1, DGS-I, nuclear factor NF90, HPIP1, and human HN1 homolog protein. Also disclosed are nucleic acids encoding the HPIP1 and human HN1 homolog protein, or derivs., fragments and analogs thereof. Methods of screening the complexes or proteins for efficacy in treating and/or

preventing certain diseases and disorders, particularly atopic diseases, autoimmune diseases, neurodegenerative disease, cancer, pigmentation disorders, platelet dysfunction and viral diseases, are also disclosed.

TC ICM C12N009-12

> ICS C07K014-47; C07K016-18; C12N015-62; C12N015-63; A61K038-17; G01N033-48

6-3 (General Biochemistry) CC

Section cross-reference(s): 1, 3, 13

Chimeric gene TΤ

RL: BUU (Biological use, unclassified); BIOL (Biological study); USES (Uses)

(for HPS-HPSIP fusion proteins; hermansky pudlak syndrome protein-interacting proteins (HPSIPs), their complexes with HPS proteins, and diagnosis and treatment of diseases)

ΙT Protein sequences

(of human proteins HPIP1 and HN1 homolog)

141933-95-1, Protein 14-3-3 (human η -chain reduced) IT 170138-42-8 175420-56-1, Atrophin 1 (human clone 1-13 gene 145110-29-8 292890-44-9 292890-45-0 292890-46-1 202012-14-4

RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL (Biological study); PROC (Process)

(amino acid sequence; hermansky pudlak syndrome protein-interacting proteins (HPSIPs), their complexes with HPS proteins, and diagnosis and treatment of diseases)

ANSWER 34 OF 86 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2000:608992 CAPLUS

DOCUMENT NUMBER: 133:187932

TITLE:

Automated, computerized toxin

screening/characterization system based on cell arrays

and fluorescent reagents

INVENTOR(S): Giuliano, Kenneth A.; Kapur, Ravi

PATENT ASSIGNEE(S): Cellomics, Inc., USA SOURCE: PCT Int. Appl., 350 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent English LANGUAGE:

FAMILY ACC. NUM. COUNT: 21

PATENT INFORMATION:

PATENT NO.										APPLICATION NO.						DATE			
WO 2000050872					A2 20000831					WO 2	000-1	JS47		20000225 <					
WO 2000050872			A3		2001	0308													
	W:	ΑE,	AL,	AM,	ΑT,	AU,	ΑZ,	BA,	BB,	ВG,	BR,	BY,	CA,	CH,	CN,	CR,	CU,		
		CZ,	DE,	DK,	DM,	EE,	ES,	FI,	GB,	GD,	GE,	GH,	GM,	HR,	HU,	ID,	IL,		
		IN,	IS,	JP,	KE,	KG,	KP,	KR,	KZ,	LC,	LK,	LR,	LS,	LT,	LU,	LV,	MA,		
		MD.	MG.	MK.	MN.	MW.	MX,	NO,	NZ,	PL,	PT,	RO,	RU,	SD,	SE,	SG,	SI,		
		-	-							UG,									
	RW:	- •	•		•	•	•	•	•	TZ,							DE,		
	2000	•	•	•			-	-	-	LU,									
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TTC	6759			•	•			•	MR, NE, SN, TD, TG US 1999-352171						19990712				
									CA 2000-2362117										
										CA 20	000-	2362.	TT/		21	0000.	225 <		
CA	2362	117			С		2004	1130											
ΑU	2000	0360	56		A5		2000	0914		AU 20	000-	3605	6		20	0000	225 <		
EР	1155	304			A2		2001	1121		EP 2	000-	9147	01		20	0000	225		
ΕP	1155	304			В1		2003	0507											
	R:	AT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR,	IT,	LI,	LU,	NL,	SE,	MC,	PT,		
		•	•	•	LV,		-	•	·	•	•	·	•	•					

AT 239907	E	20030515	ΑT	2000-914701		20000225
JP 2003526772	T2	20030909	JP	2000-601420		20000225
JP 3576491	B2	20041013				
US 2003204316	A1	20031030	US	2003-430534		20030506
US 6902883	B2	20050607				
PRIORITY APPLN. INFO.:			US	1999-122152P	P	19990226
			US	1999-123399P	P	19990308
			US	1999-352171	Α	19990712
			US	1997-810983	A2	19970227
			US	1998-31271	B2	19980227
			US	1998-92671P	P	19980713
			WO	2000-US4794	W	20000225
			US	2000-650937	A1	20000829

ED Entered STN: 01 Sep 2000

The present invention provides systems, methods, screens, reagents and AB kits for optical system anal. of cells to rapidly determine the distribution, environment, or activity of fluorescently labeled reporter mols. in cells for the purpose of screening large nos. of compds. for those that specifically affect particular biol. functions. The invention provides systems, methods, and screens that combine high throughput screening and high content screening that significantly improve target validation and candidate optimization by combining many cell screening formats with fluorescence-based mol. reagents and computer-based feature extraction, data anal., and automation, resulting in increased quantity and speed of data collection, shortened cycle times, and, ultimately, faster evaluation of promising drug candidates. For example, the effect of interleukin-1 on translocation of transcription factor NF-κB from the cytoplasm to the nucleus was analyzed using 3T3 cells in the wells of a 96-well microtiter plate. The rows of well were titered with the interleukin-1. The cells were then fixed and stained with fluorescein-labeled antibody to NF-κB and with Hoechst 33423, a DNA-specific fluorophore. Computerized fluorescent image anal. was used to compare nuclear and cytoplasm fluorescence. The decrease in this ratio was strongly correlated with concentration of interleukin-1. A number of more sophisticated assays are described.

- IC ICM G01N015-14
- CC 1-1 (Pharmacology)

Section cross-reference(s): 3

- IT Heat-shock proteins
 - RL: ARG (Analytical reagent use); ANST (Analytical study); USES (Uses) (HSP 27, fusion proteins with GFP; automated, computerized toxin screening/characterization system based on cell arrays and fluorescent reagents)
- IT Transcription factors
 - RL: ARG (Analytical reagent use); ANST (Analytical study); USES (Uses) (NF- κ B (nuclear factor κ B), **fusion** proteins with GFP; automated, computerized toxin screening/characterization system based on cell arrays and fluorescent reagents)
- IT cDNA sequences

(for GFP fusion proteins)

- IT Glucocorticoid receptors
 - Tubulins
 - RL: ARG (Analytical reagent use); ANST (Analytical study); USES (Uses) (fusion proteins with GFP; automated, computerized toxin screening/characterization system based on cell arrays and fluorescent reagents)
- IT Proteins, specific or class
 - RL: ARG (Analytical reagent use); ANST (Analytical study); USES (Uses) (green fluorescent, fusion proteins; automated, computerized toxin screening/characterization system based on cell arrays and

Agnes Rooke 10/015,956 fluorescent reagents) TT Protein sequences (of GFP fusion proteins) 289512-71-6 289512-73-8 IT 289512-69-2 RL: BUU (Biological use, unclassified); PRP (Properties); BIOL (Biological study); USES (Uses) (amino acid sequence; automated, computerized toxin screening/characterization system based on cell arrays and fluorescent reagents) 3520-43-2, JC-1 9001-92-7, Protease 23491-52-3, Hoechst 33342 IT 169592-56-7D, Caspase 3, 155215-87-5, Stress-activated protein kinase fusion protein with GFP and annexin II cytoskeletal binding domain 201860-17-5, MitoTracker Green FM 212835-04-6, Bodipy FL phallacidin RL: ARG (Analytical reagent use); ANST (Analytical study); USES (Uses) (automated, computerized toxin screening/characterization system based on cell arrays and fluorescent reagents) 143891-07-0, Phosphoprotein MAP 4 (mouse clone M31/M9/M7 IT microtubule-associated protein moiety reduced) 200014-25-1 180033-16-3 269051-21-0 269051-23-2 269051-29-8 269051-32-3 269051-19-6 269051-42-5, Calreticulin (synthetic fragment) 289514-10-9 269051-34-5 289514-62-1 289514-81-4 289514-83-6 289514-17-6 289514-15-4 289514-87-0 289514-85-8 RL: PRP (Properties) (unclaimed protein sequence; automated, computerized toxin screening/characterization system based on cell arrays and fluorescent reagents) ANSWER 35 OF 86 CAPLUS COPYRIGHT 2006 ACS on STN 2000:421333 CAPLUS ACCESSION NUMBER: 133:68897 DOCUMENT NUMBER: Yeast cell protein kinase expression system for drug TITLE: screening INVENTOR(S): Thorner, Jeremy William; Alessi, Dario Renato; Torrance, Pamela Diane; Casamayor, Antonio Medical Research Council, UK; Regents of the PATENT ASSIGNEE(S): University of California PCT Int. Appl., 154 pp. SOURCE: CODEN: PIXXD2 DOCUMENT TYPE: Patent English LANGUAGE: FAMILY ACC. NUM. COUNT: PATENT INFORMATION: APPLICATION NO. DATE DATENT NO KIND DATE

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1	WO 2000036135					A2	20000622			WO	1999-		19991214 <					
1	WO 2000036135				A3		2000	1026										
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		RW:	ΑT,	BE,	CH,	CY,	DE	, DK,	ES,	FI, FF	, GB,	GR,	ΙE,	IT,	LU,	MC,	NL,	
			PT,	SE														
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	JP	2002	5321	00		T2		2002	1002	JP	2000-	58838	33		19	9991:	214	
PRIOR	IT	APP	LN.	INFO	. :					US	1998-	11213	14P]	2 19	9981:	214	
										WO	1999-	GB422	28	1	V 19	9991:	214	
			am.		3 T	- 20	^ ^											

ED Entered STN: 23 Jun 2000

AB A method of identifying a compound which inhibits to different extents (a) a host yeast cell protein kinase or kinases and (b) a protein kinase

derivable from a source other than the said host yeast cell that is equivalent to the said host yeast cell protein kinase or kinases, wherein a compound is exposed to (1) a first host yeast cell wherein the yeast cell is capable of expressing the said host yeast cell protein kinase or kinases and is not capable of expressing the said equivalent protein kinase and (2) a second host yeast cell wherein the yeast cell is (a) not capable of expressing the said yeast cell protein kinase or kinases and (b) is capable of expressing the said equivalent protein kinase derivable from a source other than the host yeast cell and the effect of the compound on the viability of the said yeast cells is measured, and a compound that affects the viability of the first said yeast cell and the said second yeast cell differently, is identified. The method may be useful in a screen for identifying compds. that inhibit a mammalian or fungal protein kinase. The compds. may be useful in medicine.

IC ICM C12Q001-00

CC 1-1 (Pharmacology)

Section cross-reference(s): 3

IT Antitumor agents

Aspergillus

Aspergillus flavus

Aspergillus fumigatus

Aspergillus niger

Blastomyces

Blastomyces dermatitidis

Botryoascus

Candida

Candida albicans

Cephalosporium

Citeromyces

Cladosporium carrionii

Coccidioides

Coccidioides immitis

Cryptococcus (fungus)

Cryptococcus neoformans

Debaryomyces

Drug screening

Endomycopsis

Epidermophyton

Fonsecaea pedrosoi

Fungicides

Fusarium

Hansenula

Histoplasma

Histoplasma capsulatum

Kluyveromyces

Leucosporidium

Madurella

Madurella grisea

Madurella mycetomatis

Malassezia

Malassezia furfur

Metschnikowia

Microsporum

Molecular cloning

Pachysolen (fungus)

Paracoccidioides

Paracoccidioides brasiliensis

Phialophora compacta

Phialophora verrucosa

Phosphorylation, biological

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Pichia
     Pneumocystis carinii
       Protein sequences
     Prototheca
     Prototheca wickerhamii
     Pseudallescheria boydii
     Rhinocladiella aquaspersa
     Rhizomucor
     Rhizopus
     Rhodosporidium
     Saccharomyces
     Saccharomyces cerevisiae
     Schizosaccharomyces
     Sporidiobolus
     Sporothrix
     Sporothrix schenckii
     Test kits
     Torulopsis
     Trichophyton
     Yeast
     cDNA sequences
        (yeast cell protein kinase expression system for drug screening)
     Fusion proteins (chimeric proteins)
     RL: BPN (Biosynthetic preparation); THU (Therapeutic use); BIOL
     (Biological study); PREP (Preparation); USES (Uses)
        (yeast cell protein kinase expression system for drug screening)
     124671-35-8
                   225792-13-2
                                 277762-90-0
                                                277762-92-2
     277762-94-4
                   277763-15-2
                                  278189-91-6
                                                278189-92-7
                                                              278189-93-8
     278189-94-9
                   278189-95-0
                                  278189-96-1
                                                278779-52-5
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                                  278779-56-9
                                                278779-57-0
                                                              278779-58-1
     278779-54-7
     278779-59-2
     RL: PRP (Properties)
        (unclaimed protein sequence; yeast cell protein
        kinase expression system for drug screening)
     ANSWER 36 OF 86 CAPLUS COPYRIGHT 2006 ACS on STN
                         2000:402007 CAPLUS
ACCESSION NUMBER:
DOCUMENT NUMBER:
                         133:53686
TITLE:
                         Chlamydial antigens and genomic DNA sequences for
                         treatment and diagnosis of chlamydial infection
INVENTOR(S):
                         Probst, Peter; Bhatia, Ajay; Skeiky, Yasir A. W.;
                         Fling, Steven P.; Jen, Shyian; Stromberg, Erica Jean
PATENT ASSIGNEE(S):
                         Corixa Corporation, USA
SOURCE:
                         PCT Int. Appl., 256 pp.
                         CODEN: PIXXD2
DOCUMENT TYPE:
                         Patent
                         English
LANGUAGE:
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
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WO 2000	0344	83		A2		2000	0615	1	WO 1	999-1	US29	012		19	99912	208 <	-
WO 2000	0344	83		A3	:	2001	1101										
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	MD,	MG,	MK,	MN,	MW,	MX,	NO,	NZ,	PL,	PT,	RO,	RU,	SD,	SE,	SG,	SI,	
	SK,	SL,	ТJ,	TM,	TR,	TT,	TZ,	UA,	UG,	US,	UZ,	VN,	ΥU,	ZA,	ZW,	AM,	
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IT

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RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE,
             DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF,
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     EP 1144642
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                                            EP 1999-963037
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     EP 1144642
                          A3
                                20020605
            AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, SI, LT, LV, FI, RO
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                                20031219
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     AU 769293
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     NO 2001002812
                         Α
                                20010802
                                            NO 2001-2812
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PRIORITY APPLN. INFO.:
                                            US 1998-208277
                                                                A 19981208
                                            US 1999-288594
                                                                A 19990408
                                            US 1999-410568
                                                                A 19991001
                                            US 1999-426571
                                                                A 19991022
                                            WO 1999-US29012
                                                                   19991208
    Entered STN: 16 Jun 2000
ED
AB
     Compds. and methods for the diagnosis and treatment of Chlamydial
     infection are disclosed. The compds. provided include polypeptides that
     contain at least one antigenic portion of a Chlamydia antigen and DNA
     sequences encoding such polypeptides. Chlamydia antigens were isolated by
     expression cloning of a genomic DNA library of C. trachomatis LGV II, and
     shown to induce T cell proliferation and interferon-β production Immune
     responses of human PBMC and T cell lines are generated against the
     Chlamydia antigens. Pharmaceutical compns. and vaccines comprising such
    polypeptides or DNA sequences are also provided, together with antibodies
    directed against such polypeptides. Diagnostic kits containing such
    polypeptides or DNA sequences and a suitable detection reagent may be used
     for the detection of Chlamydial infection in patients and in biol.
     samples.
IC
     ICM C12N015-31
         C07K014-295; C12N001-21; C12N001-19; C12N005-10; C07K019-00;
     ICS
          C12N015-62; C07K016-02; A61K039-118; A61K031-70; G01N033-569;
          C12Q001-68; A61K048-00
CC
     1-5 (Pharmacology)
     Section cross-reference(s): 3, 10, 15
IT
    Chlamydia
    Chlamydia pneumoniae
    Chlamydia trachomatis
    DNA sequences
    Molecular cloning
      Protein sequences
     Vaccines
        (Chlamydial antigens and genomic DNA sequences for treatment and
        diagnosis of chlamydial infection)
IT
    Fusion proteins (chimeric proteins)
    RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (Chlamydial antigens and genomic DNA sequences for treatment and
        diagnosis of chlamydial infection)
IT
    223705-51-9
                  261940-10-7
                                 275830-86-9
                                               275830-88-1
                                                             275830-92-7
    275830-94-9
                  275830-96-1
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                                               276254-68-3
                                                             276254-72-9
    275831-25-9
    RL: PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES
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(Uses) (amino acid sequence; Chlamydial antigens and genomic DNA sequences for treatment and diagnosis of chlamydial infection) 275830-76-7 275830-77-8 275830-78-9 IT 275830-75-6 275830-79-0 275830-83-6 275830-80-3 275830-81-4 275830-82-5 275830-84-7 275830-85-8 275830-87-0 275830-89-2 275830-90-5 275830-91-6 275830-95-0 275830-97-2 275830-98-3, DNA (Chlamydia 275830-93-8 trachomatis gene pmpI) 275830-99-4, DNA (Chlamydia trachomatis gene 275831-00-0, DNA (Chlamydia trachomatis gene pmpE) 275831-01-1, DNA (Chlamydia trachomatis gene pmpD) 275831-02-2, DNA (Chlamydia trachomatis gene pmpC) 275831-03-3, DNA (Chlamydia trachomatis gene pmpB) 275831-04-4, DNA (Chlamydia trachomatis gene pmpI) 275831-11-3, DNA (Chlamydia trachomatis gene pmpG) 275831-10-2 275831-12-4 275831-13-5 275831-14-6 275831-15-7 275831-16-8 275831-17-9 275831-26-0 275831-28-2 275831-30-6 275831-31-7 275831-32-8 275831-33-9 275831-34-0 275831-35-1 275831-36-2 275831-37-3 275831-38-4 275831-39-5 275831-40-8 275831-41-9 275831-42-0 275831-43-1 275831-44-2 275831-45-3 275831-46-4 275831-47-5 275831-48-6 275831-49-7 275831-50-0 275831-51-1 275831-52-2 275831-53-3, DNA (Chlamydia pneumoniae gene CT529) RL: PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (nucleotide sequence; Chlamydial antigens and genomic DNA sequences for treatment and diagnosis of chlamydial infection) IT 275832-48-9 275832-49-0 275832-50-3 275832-51-4 275832-52-5 275832-53-6 275832-54-7 275832-55-8 275832-56-9 275832-57-0 RL: PRP (Properties) (unclaimed protein sequence; chlamydial antigens and genomic DNA sequences for treatment and diagnosis of chlamydial infection) ANSWER 37 OF 86 CAPLUS COPYRIGHT 2006 ACS on STN 2000:384389 CAPLUS ACCESSION NUMBER: DOCUMENT NUMBER: 133:27383 Plant vitamin E biosynthetic enzymes TITLE: γ-tocopherol methyltransferase and 4-hydroxyphenylpyruvate dioxygenase and cDNAs and their uses INVENTOR(S): Cahoon, Rebecca E.; Coughlan, Sean J.; Miao, Guo-hua; Rafalski, J. Antoni E. I. Du Pont de Nemours & Co., USA PATENT ASSIGNEE(S): SOURCE: PCT Int. Appl., 82 pp. CODEN: PIXXD2 DOCUMENT TYPE: Patent LANGUAGE: English FAMILY ACC. NUM. COUNT: PATENT INFORMATION: PATENT NO. KIND DATE APPLICATION NO. DATE ------------------------WO 2000032757 A2 20000608 WO 1999-US28588 19991202 <--**A3** WO 2000032757 20001123 AE, AL, AU, BA, BB, BG, BR, CA, CN, CR, CU, CZ, DM, EE, GD, GE, HR, HU, ID, IL, IN, IS, JP, KP, KR, LC, LK, LR, LT, LV, MG, MK, MN, MX, NO, NZ, PL, RO, SG, SI, SK, SL, TR, TT, UA, US, UZ, VN, YU, ZA, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG 20050901

A1

RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF,

US 2005-51785

20050204

US 2005193445

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US 2005216973
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                                20050929
                                            US 2005-141593
                                                                   20050531
PRIORITY APPLN. INFO.:
                                            US 1998-110781P
                                                                P 19981203
                                            WO 1999-US28588
                                                                W 19991202
                                            US 2001-857613
                                                                A3 20010604
ED
     Entered STN: 09 Jun 2000
     This invention relates to isolated cDNA fragments encoding vitamin E
AB
     biosynthetic enzyme \gamma-tocopherol methyltransferase from corn, rice,
     soybean, and wheat and 4-hydroxyphenylpyruvate dioxygenase from rice,
     soybean, wheat, Vernonia, and Catalpa. The invention also relates to the
     construction of a chimeric gene encoding all or a portion of the vitamin E
     biosynthetic enzyme, in sense or antisense orientation, wherein expression
     of the chimeric gene results in production of altered levels of the vitamin E
     biosynthetic enzyme in a transformed host cell. The cDNAs may be used in
     selection of transformed cells as well as for screening for vitamin
     E-biosynthesis enzyme-inhibiting herbicides.
IC
     ICM C12N009-00
     3-3 (Biochemical Genetics)
CC
     Section cross-reference(s): 7, 11
     Catalpa
IT
     Corn
       Protein sequences
     Rice (Oryza sativa)
     Soybean (Glycine max)
     Vernonia mespilifolia
     Wheat
     cDNA sequences
        (plant vitamin E biosynthetic enzymes \gamma-tocopherol
        methyltransferase and 4-hydroxyphenylpyruvate dioxygenase and cDNAs and
        their uses)
IT
     Chimeric gene
     RL: AGR (Agricultural use); ARG (Analytical reagent use); BUU (Biological
     use, unclassified); ANST (Analytical study); BIOL (Biological study); USES
        (plant vitamin E biosynthetic enzymes \gamma-tocopherol
        methyltransferase and 4-hydroxyphenylpyruvate dioxygenase and cDNAs and
        their uses)
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                                                             273718-47-1
     RL: AGR (Agricultural use); BAC (Biological activity or effector, except
     adverse); BSU (Biological study, unclassified); PRP (Properties); BIOL
     (Biological study); USES (Uses)
        (amino acid sequence; plant vitamin E biosynthetic enzymes
        \gamma-tocopherol methyltransferase and 4-hydroxyphenylpyruvate
       dioxygenase and cDNAs and their uses)
                  194615-50-4
IT
     170977-25-0
                                 195397-27-4
                                               219903-83-0
                                                             220379-64-6
     RL: PRP (Properties)
        (unclaimed protein sequence; plant vitamin E
        biosynthetic enzymes \gamma-tocopherol methyltransferase and
        4-hydroxyphenylpyruvate dioxygenase and cDNAs and their uses)
    ANSWER 38 OF 86 CAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                         2000:314815 CAPLUS
DOCUMENT NUMBER:
                         132:344102
TITLE:
                         Gene cluster encoding oleandolide synthases from
                         Streptomyces antibioticus and recombinant expression
                         of hybrid genes encoding heterologous polyketide
                         synthases
INVENTOR (S):
                         Betlach, Mary C.; Shah, Sanjay Krishnakant; Mcdaniel,
```

Robert; Tang, Li

PATENT ASSIGNEE(S):

Kosan Biosciences, Inc., USA

SOURCE:

PCT Int. Appl., 86 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PA	rent n	ю.			KIN	D D	ATE		AP	PLICAT	ION	NO.		I	ATE		
	20000							0511 0831	WO	1999-	US24	478		1	9991	022	<
	W: RW:	•	BE,		CY,	DE,	DK,	ES,	FI, F	R, GB,	GR,	IE,	IT,	LU,	MC,	NL,	•
CA	23474	12			AA	2	000	0511	CA	1999-	2347	412		1	19991	022	<
EP	11249	69			A2	2	001	0822	EP	1999-	9550	63		1	19991	022	
	R:	AT, IE,		CH,	DE,	DK,	ES,	FR,	GB, G	R, IT,	LI,	LU,	NL,	SE	MC,	PT,	•
AU	75842	1			B2	2	003	0320	ΑŪ	2000-	1125	4]	19991	022	
JP	20035						003	0402	JP	2000-	5797	21		1	L9991	022	
	62516				B1	2	001	0626	US	1999-	4285	17		1	19991	028	
US	20010	340	46		A1	2	001	1025	US	2001-	7689	27		2	20010	123	
ບຣ	63880	99			B2	2	002	0514									
US	20030	272	87		A1	2	003	0206	US	2001-	8088	80		2	20010	314	
US	20021	.687	30		A1	2	002	1114	US	2001-	9914	48		2	20011	116	
US	68281	.26			B2	2	004	1207									
PRIORIT	Y APPI	.N.	INFO	.:					US	1998-	1061	00P]	P 1	L9981	029	
									US	1999-	1202	54P]	P :	L9990	216	
									WO	1999-	US24	478	Ţ	N :	19991	022	
									US	1999-	4285	17	i	A1 1	19991	028	
									US	2000-	1776	60P]	P 2	20000	127	
									US	2001-	7689	27	Ī	A3 2	20010	123	

Entered STN: 15 May 2000 ED

(of oleandolide synthases from Streptomyces antibioticus)

Recombinant DNA compds. that encode all or a portion of the oleandolide AB polyketide synthase are used to express recombinant polyketide synthase genes in host cells for the production of oleandolide, oleandolide derivs., and polyketides that are useful as antibiotics and motilides. Genomic DNA containing the complete 8,8a-deoxyoleandrolide synthase gene cluster isolated from an oleandomycin-reproducing strain of Streptomyces antibioticus comprises 3 synthase open reading frames (oleAI, oleAII, and oleAII), each ORF comprising 2 extender modules and the first ORF also encoding the loading module, plus a number of tailoring enzyme genes. Each ORF is composed of at least a ketosynthase, an acyltransferase, and an acyl carrier protein domain. Hybrid polyketide synthase enzymes are prepared by fusion of the loading module of either 5-deoxyerythronolide B synthase or narbonolide synthase with extender modules 1 and 2 of the oleandolide synthase (gene oleAI) and coexpressed with oleAII and oleAIII genes in suitable host cells such as Streptomyces lividans or Saccharopolyspora erythraea. Another example of a hybrid polyketide synthase is prepared by co-expressing the oleAI and oleAII genes with an oleAIII gene encoding extender module 5 and the ketosynthase and acyltransferase of extender module 6 of the oleandolide synthase fused to the acyl carrier protein of extender module 6 and the thioesterase of narbonolide synthase.

ICM C12N009-00 IC

^{3-2 (}Biochemical Genetics) CC

Section cross-reference(s): 1, 7, 10, 16

Protein sequences IT

IT268532-04-3P 268532-05-4P **268532-06-5P** RL: BPN (Biosynthetic preparation); BUU (Biological use, unclassified); CAT (Catalyst use); PRP (Properties); BIOL (Biological study); PREP (Preparation); USES (Uses) (amino acid sequence; gene cluster encoding oleandolide synthases from Streptomyces antibioticus and recombinant expression of hybrid genes encoding heterologous polyketide synthases) 79956-01-7DP, Polyketide synthase, fusion proteins with IT oleandolide synthase 128172-72-5DP, 6-Deoxyerythronolide B synthase, fusion proteins with oleandolide synthase 247595-90-0DP, Picromycin synthase, fusion proteins with oleandolide synthase 251091-63-1DP, Narbonolide synthase, fusion proteins with oleandolide synthase RL: BPN (Biosynthetic preparation); CAT (Catalyst use); BIOL (Biological study); PREP (Preparation); USES (Uses) (gene cluster encoding oleandolide synthases from Streptomyces antibioticus and recombinant expression of hybrid genes encoding heterologous polyketide synthases) ANSWER 39 OF 86 CAPLUS COPYRIGHT 2006 ACS on STN 2000:278133 CAPLUS ACCESSION NUMBER: DOCUMENT NUMBER: 132:319509 TITLE: An improved method for extracting quantitative information relating to an influence on a cellular response Arkhammar, Per O. G.; Terry, Bernard Robert; Scudder, INVENTOR (S): Kurt Marshall; Bjorn, Sara Petersen; Thastrup, Ole; Hagel, Grith PATENT ASSIGNEE(S): Bioimage A/S, Den. PCT Int. Appl., 150 pp. SOURCE: CODEN: PIXXD2 DOCUMENT TYPE: Patent English LANGUAGE: FAMILY ACC. NUM. COUNT: PATENT INFORMATION: APPLICATION NO. DATE PATENT NO. KIND DATE --------------------WO 2000023615 A2 WO 2000023615 A3 WO 1999-DK562 20000427 19991015 <--20010329 W: AE, AL, AM, AT, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, CZ, DE, DE, DK, DK, DM, EE, EE, ES, FI, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG 20000508 AU 1999-61895 AU 9961895 A1 19991015 <--19991015 20010808 EP 1999-948731 EP 1121593 A2 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO T2 JP 2002527761 20020827 JP 2000-577322 19991015 DK 1998-1320 PRIORITY APPLN. INFO.: A 19981015

ED Entered STN: 28 Apr 2000

AB An improved method and tools for quantifying the effect of an influence on cellular response is described. In particular, an improved method is described for detecting intracellular translocation or redistribution of

WO 1999-DK562

W 19991015

biol. active polypeptides. The invention also describes several ways of contacting the cells with a substance influencing a cellular response and extracting quant. information relating to the response in a highly parallel fashion. The method may be used as a very efficient procedure for testing or discovering the influence of a substance on a physiol. process using com. available parallel, high volume assay techniques, for example in connection with screening for new drugs, testing of substances for toxicity, and identifying drug targets for known or novel drugs. A fusion protein having the catalytic subunit of mouse protein kinase A fused with [64-leucine, 65-threonine] green fluorescent protein was prepared by recombinant methods. To assess the effect of glucagon on fusion protein translocation, the fusion protein was stably expressed in BHK cells overexpressing the human glucagon receptor. ICM C12Q001-68 9-5 (Biochemical Methods) Section cross-reference(s): 1, 3, 4, 6 cell response assay; translocation fusion protein green fluorescent protein; kinase fusion green fluorescent protein translocation; biol signal transduction Transcription factors RL: ARG (Analytical reagent use); BPN (Biosynthetic preparation); BPR (Biological process); BSU (Biological study, unclassified); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); PREP (Preparation); PROC (Process); USES (Uses) (NF-κB (nuclear factor κB), fusion proteins with enhanced green fluorescent protein; improved method for extracting quant. information relating to an influence on a cellular response) Transcription factors RL: ARG (Analytical reagent use); BPN (Biosynthetic preparation); BPR (Biological process); BSU (Biological study, unclassified); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); PREP (Preparation); PROC (Process); USES (Uses) (Smad-2, fusion proteins with enhanced green fluorescent protein; improved method for extracting quant. information relating to an influence on a cellular response) Phosphoproteins RL: ARG (Analytical reagent use); BPN (Biosynthetic preparation); BPR (Biological process); BSU (Biological study, unclassified); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); PREP (Preparation); PROC (Process); USES (Uses) (VASP (vasodilator-stimulated protein), fusion proteins with enhanced green fluorescent protein; improved method for extracting quant. information relating to an influence on a cellular response) Chimeric gene RL: ARG (Analytical reagent use); BPN (Biosynthetic preparation); BPR (Biological process); BSU (Biological study, unclassified); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); PREP (Preparation); PROC (Process); USES (Uses) (for fluorophore, expression of, in cell; improved method for extracting quant. information relating to an influence on a cellular response) DNA sequences Drug screening Fluorometry Imaging Microtiter plates Molecular cloning Protein sequences

Signal transduction, biological

IC

CC

ST

IT

TT

TT

TT

Transformation, genetic Video cameras (improved method for extracting quant. information relating to an influence on a cellular response) IT Fusion proteins (chimeric proteins) RL: ARG (Analytical reagent use); BPN (Biosynthetic preparation); BPR (Biological process); BSU (Biological study, unclassified); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); PREP (Preparation); PROC (Process); USES (Uses) (with green fluorescent protein; improved method for extracting quant. information relating to an influence on a cellular response) TT 214766-59-3P 214767-46-1P 214768-64-6P 214768-70-4P 214768-72-6P 214769-08-1P 214769-33-2P 214769-36-5P 266330-89-6P RL: ARG (Analytical reagent use); BPN (Biosynthetic preparation); BPR (Biological process); BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); PREP (Preparation); PROC (Process); USES (Uses) (amino acid sequence; improved method for extracting quant. information relating to an influence on a cellular response) 142008-29-5P, Protein kinase A IT RL: ARG (Analytical reagent use); BPN (Biosynthetic preparation); BPR (Biological process); BSU (Biological study, unclassified); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); PREP (Preparation); PROC (Process); USES (Uses) (catalytic fragment, fusion proteins with green fluorescent protein; improved method for extracting quant. information relating to an influence on a cellular response) IT 9007-92-5, Glucagon, biological studies RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL (Biological study); PROC (Process) (fluorescent protein kinase C fusion protein translocation response to; improved method for extracting quant. information relating to an influence on a cellular response) IT 137632-07-6P, Kinase (phosphorylating), protein, ERK1 RL: ARG (Analytical reagent use); BPN (Biosynthetic preparation); BPR (Biological process); BSU (Biological study, unclassified); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); PREP (Preparation); PROC (Process); USES (Uses) (fusion proteins with green fluorescent protein; improved method for extracting quant. information relating to an influence on a cellular response) IT 141436-78-4P RL: ARG (Analytical reagent use); BPN (Biosynthetic preparation); BPR (Biological process); BSU (Biological study, unclassified); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); PREP (Preparation); PROC (Process); USES (Uses) $(\alpha, \alpha \text{ and } \beta 1,$ fusion proteins with green fluorescent protein; improved method for extracting quant. information relating to an influence on a cellular response) ANSWER 40 OF 86 CAPLUS COPYRIGHT 2006 ACS on STN ACCESSION NUMBER: 2000:277863 CAPLUS DOCUMENT NUMBER: 132:303473 TITLE: Specific therapeutic interventions obtained by interference with redistribution and/or targetting of cyclic nucleotide phosphodiesterases or Ik kinases INVENTOR (S): Arkhammar, Per O. G.; Terry, Bernard Robert; Scudder, Kurt Marshall; Bjorn, Sara Petersen; Thastrup, Ole PATENT ASSIGNEE(S): Bioimage A/s, Den.

PCT Int. Appl., 128 pp. SOURCE:

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

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PATENT NO.
                               KIND
                                        DATE
                                                      APPLICATION NO.
                                                       ______
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                               _ _ _ _
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      WO 2000023091
                                A2
                                        20000427
                                                       WO 1999-DK567
                                                                                    19991015 <--
                                        20000713
      WO 2000023091
                               A3
               AE, AL, AM, AT, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, CZ, DE, DE, DK, DK, DM, EE, EE, ES, FI, FI, GB, GD, GE,
                GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT,
                RO, RU, SD, SE, SG, SI, SK, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
          RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF,
                CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
                                                    CA 1999-2346937
                                        20000427
                                                                                    19991015 <--
      CA 2346937
                                AA
                                                    AU 1999-61899
EP 1999-948735
      AU 9961899
                                A1
                                        20000508
                                                                                    19991015 <--
                                        20011024
                                                                                    19991015
                                A2
      EP 1146888
                AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
                IE, SI, LT, LV, FI, RO
                                                       JP 2000-576864
      JP 2003526608
                               T2
                                        20030909
                                                                                     19991015
                                                       DK 1998-1321
PRIORITY APPLN. INFO.:
                                                                                A 19981015
                                                       DK 1998-1322
                                                                                A 19981015
                                                       DK 1998-1323
                                                                                A 19981015
                                                       WO 1999-DK567
                                                                                W 19991015
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Entered STN: 28 Apr 2000 ED

- The application describes a novel mechanism of action, that is modulation AΒ of the specific effectiveness of I-kappa-kinases or cyclic nucleotide phosphodiesterases (PDEs) which have the ability to cleave cGMP or cAMP. The preferred mode of action is dislocation, disruption of targeting or interference with redistribution of specific isoforms or splice variants of PDE4, PDE5, or I-kappa-kinases from their anchoring sites within cells, thereby modulating their specific effectiveness, not their enzymic capacity. The chemical entities may be useful in preventing or treating in an animal, preferably a human, in need thereof an adverse condition which may be reduced or abolished by modulating the specific effectiveness of PDE4, PDE5, or I-kappa-kinases.
- IC ICM A61K038-00
- CC 1-1 (Pharmacology)

Section cross-reference(s): 3

IT Aequorea victoria

Anti-inflammatory agents

Antiasthmatics

Antidepressants

Antidiabetic agents

Antihypertensives

Antihypotensives

Antirheumatic agents

Asthma

Autoimmune disease

Drug screening

Fluorescent probes

Fluorometry

Graves' disease

Hypertension

Hypotension
Inflammation
Molecular cloning
Myasthenia gravis
Protein sequences
Rheumatoid arthritis

Rifeumatoru artifrit.

cDNA sequences

(specific therapeutic interventions obtained by interference with redistribution and/or targetting of cyclic nucleotide phosphodiesterases or $I\kappa$ kinases)

IT Fusion proteins (chimeric proteins)

RL: BOC (Biological occurrence); BPR (Biological process); BSU (Biological study, unclassified); PRP (Properties); BIOL (Biological study); OCCU (Occurrence); PROC (Process)

(specific therapeutic interventions obtained by interference with redistribution and/or targetting of cyclic nucleotide phosphodiesterases or $I\kappa$ kinases)

IT 265295-40-7P 265295-42-9P 265295-44-1P 265295-46-3P 265295-48-5P **265295-51-0P** 265295-53-2P 265295-55-4P

RL: BAC (Biological activity or effector, except adverse); BOC (Biological occurrence); BPN (Biosynthetic preparation); BPR (Biological process); BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); OCCU (Occurrence); PREP (Preparation); PROC (Process); USES (Uses)

(amino acid sequence; specific therapeutic interventions obtained by interference with redistribution and/or targetting of cyclic nucleotide phosphodiesterases or Ik kinases)

L8 ANSWER 41 OF 86 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 200

2000:220716 CAPLUS

DOCUMENT NUMBER:

132:261375

TITLE:

Immunoglobulin fusion product with

immunoglobulin receptor that protects Ig in mucosal environment, cDNA sequences, transgenic plants, and

dental caries prevention

INVENTOR(S):
Hiatt, Andrew C.; Ma, Julian K. C.; Lehner, Thomas;

Mostov, Keith E.

PATENT ASSIGNEE(S): USA

SOURCE: U.S., 59 pp., Cont.-in-part of U.S. Ser. No. 367,395.

CODEN: USXXAM

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

PATENT NO			KIN	D DATE		APPLICATION NO. DATE	
US 604603	7		Α	20000	0404	US 1995-434000 19950504 <	-
CA 220878	3		AA	19960	711	CA 1995-2208783 19951227 <	-
WO 962101	2		A1	19960	0711	WO 1995-US16889 19951227 <	-
W: A	J, BR,	CA,	CN,	CZ, FI,	HU,	JP, KR, MX, NO, NZ, PL, RU, SG	
RW: A	C, BE,	CH,	DE,	DK, ES,	FR,	GB, GR, IE, IT, LU, MC, NL, PT, SE	
AU 964608	3		A1	19960	724	AU 1996-46088 19951227 <	-
AU 722668			В2	20000	0810		
EP 807173			A 1	19971	1119	EP 1995-944237 19951227 <	-
R: A	, BE,	CH,	DE,	DK, ES,	FR,	GB, GR, IT, LI, LU, NL, SE, MC, PT, IE	3
CN 118380	?		A	19980	0603	CN 1995-197699 19951227 <	-
US 630334	L		B1	20011	1016	US 1999-312157 19990514	
AU 773602			B2	20040	0527	AU 2000-71534 20001110	
US 680870)		B1	20041	1026	US 2000-717888 20001120	

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US 2002159958
                                            US 2001-982107
                                                                   20011016
                         A1
                                20021031
                                            US 2004-781989
                                                                   20040218
     US 2005202026
                         A1
                                20050915
PRIORITY APPLN. INFO.:
                                            US 1994-367395
                                                                B2 19941230
                                            US 1995-434000
                                                                A 19950504
                                            AU 1996-46088
                                                                A3 19951227
                                            WO 1995-US16889
                                                                W 19951227
                                            US 1999-312157
                                                                A1 19990514
                                            US 2000-717888
                                                                A1 20001120
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ED Entered STN: 06 Apr 2000

AB Igs of the present invention are useful as therapeutic Igs against mucosal pathogens such as Streptococcus mutans. The Igs contain a protection protein (e.g., the polyimmunoglobulin receptor) that protects the Igs in the mucosal environment. The invention also includes a greatly improved method of producing Igs in plants by producing the protection protein in the same cell as the other components of the Igs. The components of the Ig are assembled at a much improved efficiency. The method of the invention allows the assembly and high efficiency production of such complex mols. The invention also contemplates the production of Igs containing protection

proteins in a variety of cells, including plant cells, that can be selected for useful addnl. properties. The use of Igs containing protection proteins as therapeutic antibodies against mucosal and other pathogens is also contemplated.

IC ICM C12N015-00

ICS C12N015-29; C12N015-82; A01H004-00

INCL 435070100

CC 3-2 (Biochemical Genetics)

Section cross-reference(s): 1, 11, 15

- ST Ig fusion receptor protection mucosa caries; dental caries prevention Ig fusion receptor; sequence Ig fusion receptor mucosa protection; plant transgenic manuf Ig fusion receptor
- IT Immunoglobulins

RL: BMF (Bioindustrial manufacture); BPN (Biosynthetic preparation); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(A; Ig **fusion** product with Ig receptor that protects Ig in mucosal environment, cDNA sequences, transgenic plants, and dental caries prevention)

IT Immunoglobulins

RL: BMF (Bioindustrial manufacture); BPN (Biosynthetic preparation); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(D; Ig fusion product with Ig receptor that protects Ig in mucosal environment, cDNA sequences, transgenic plants, and dental caries prevention)

IT Immunoglobulins

RL: BMF (Bioindustrial manufacture); BPN (Biosynthetic preparation); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(E; Ig fusion product with Ig receptor that protects Ig in mucosal environment, cDNA sequences, transgenic plants, and dental caries prevention)

IT Immunoglobulins

RL: BMF (Bioindustrial manufacture); BPN (Biosynthetic preparation); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(G; Ig fusion product with Ig receptor that protects Ig in mucosal environment, cDNA sequences, transgenic plants, and dental caries prevention)

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IT
     Immunoglobulins
     RL: BMF (Bioindustrial manufacture); BPN (Biosynthetic preparation); PRP
     (Properties); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (Guy's 13, fusion products, with Ig receptors; Ig
        fusion product with Iq receptor that protects Iq in mucosal
        environment, cDNA sequences, transgenic plants, and dental caries
        prevention)
TТ
     Antigens
    RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (Ig antigen-binding domain; Ig fusion product with Ig
        receptor that protects Ig in mucosal environment, cDNA sequences,
        transgenic plants, and dental caries prevention)
     Agrobacterium tumefaciens
IT
     Alfalfa (Medicago sativa)
     Arabidopsis
    Dicotyledon (Magnoliopsida)
     Immunotherapy
    Monocotyledon (Liliopsida)
    Mucous membrane
     Petunia
      Protein sequences
     Streptococcus mutans
     Streptococcus sobrinus
     Tobacco
    Tomato
     cDNA sequences
        (Ig fusion product with Ig receptor that protects Ig in
        mucosal environment, cDNA sequences, transgenic plants, and dental
        caries prevention)
IT
     Immunoglobulins
    RL: BMF (Bioindustrial manufacture); BPN (Biosynthetic preparation); PRP
     (Properties); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (M; Ig fusion product with Ig receptor that protects Ig in
        mucosal environment, cDNA sequences, transgenic plants, and dental
        caries prevention)
IT
    Tooth
        (caries, prevention of; Ig fusion product with Ig receptor
        that protects Ig in mucosal environment, cDNA sequences, transgenic
       plants, and dental caries prevention)
IT
     Immunoglobulin receptors
    RL: BMF (Bioindustrial manufacture); BPN (Biosynthetic preparation); PRP
     (Properties); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (fusion products, with Igs; Ig fusion product with
        Ig receptor that protects Ig in mucosal environment, cDNA sequences,
        transgenic plants, and dental caries prevention)
IT
    Transformation, genetic
        (transgenic, Ig fusion product with Ig receptor that protects
        Ig in mucosal environment, cDNA sequences, transgenic plants, and
        dental caries prevention)
IT
    144997-23-9DP, Glycoprotein (human secretory component protein moiety
    reduced), fusion products with Ig 170979-93-8DP,
     fusion products with Ig
                              180616-69-7DP, Receptor, immunoglobulin
     (rabbit), fusion products with Ig 180616-70-0DP,
    Receptor, immunoglobulin (mouse), fusion products with Ig
    180686-83-3DP, Receptor, immunoglobulin (rat), fusion
    products with Ig 180686-85-5DP, fusion products with Ig
    receptor 180686-87-7DP, fusion products with Ig receptor
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RL: BPN (Biosynthetic preparation); PRP (Properties); THU (Therapeutic
    use); BIOL (Biological study); PREP (Preparation); USES (Uses)
        (amino acid sequence; Ig fusion product with Ig receptor that
       protects Ig in mucosal environment, cDNA sequences, transgenic plants,
       and dental caries prevention)
                                           140262-61-9DP,
    140080-71-3DP, fusion products with Ig
IT
    fusion products with Ig
                              153420-82-7DP, fusion products
              153665-28-2DP, fusion products with Ig
    with Ig
    159070-18-5DP, fusion products with Ig 180686-84-4DP,
    fusion products with Ig receptor cDNA 180686-86-6DP,
    fusion products with Ig receptor cDNA
    RL: BPN (Biosynthetic preparation); PRP (Properties); THU (Therapeutic
    use); BIOL (Biological study); PREP (Preparation); USES (Uses)
        (nucleotide sequence; Ig fusion product with Ig receptor that
       protects Iq in mucosal environment, cDNA sequences, transgenic plants,
       and dental caries prevention)
    244135-92-0, PN: US5959177 SEQID: 13 unclaimed DNA
                                                         244135-94-2, PN:
    US5959177 SEQID: 14 unclaimed DNA 244135-95-3, PN: US5959177 SEQID: 15
                   244135-96-4, PN: US5959177 SEQID: 16 unclaimed DNA
    unclaimed DNA
    244135-97-5, PN: US5959177 SEQID: 17 unclaimed DNA
    RL: PRP (Properties)
        (unclaimed nucleotide sequence; Ig fusion product with Ig
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REFERENCE COUNT:
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    ANSWER 42 OF 86 CAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
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DOCUMENT NUMBER:
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                        Recombinant antigen immunoassay for the diagnosis of
TITLE:
                        syphilis
                        Mullenix, Michael C.; Deutsch, John
INVENTOR(S):
PATENT ASSIGNEE(S):
                        Becton, Dickinson and Company, USA
                        Eur. Pat. Appl., 16 pp.
SOURCE:
                        CODEN: EPXXDW
DOCUMENT TYPE:
                        Patent
LANGUAGE:
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FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
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PRIORITY APPLN. INFO.:
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    A method for detecting anti-Treponema pallidum antibody and diagnosing
AB
    syphilis has been provided. Fusion protein antigens from the fusion of a
    peptide sequence having an amino acid sequence encoded by the described
    nucleic acid sequence to Treponema pallidum membrane proteins are used as
    antigens in immunoassay of test samples for the presence of anti-Treponema
    pallidum membrane protein antibodies. A test kit for diagnosing syphilis
    is also provided comprising a container having therein the fusion protein
    antigens.
    ICM G01N033-571
IC
     ICS G01N033-543; G01N033-558; C12N015-30; C07K014-44
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CC 9-10 (Biochemical Methods) Section cross-reference(s): 3, 14, 15 IT Blood analysis DNA sequences Diagnosis Protein sequences Syphilis Treponema pallidum (recombinant antigen immunoassay for diagnosis of syphilis) 261151-34-2 261151-36-4 261151-38-6 IT RL: BUU (Biological use, unclassified); PRP (Properties); BIOL (Biological study); USES (Uses) (amino acid sequence; recombinant antigen immunoassay for diagnosis of syphilis) IT 58-85-5 RL: ARG (Analytical reagent use); BPR (Biological process); BSU (Biological study, unclassified); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); PROC (Process); USES (Uses) (fusion with recombinant antiqens; recombinant antiqen immunoassay for diagnosis of syphilis) ANSWER 43 OF 86 CAPLUS COPYRIGHT 2006 ACS on STN ACCESSION NUMBER: 2000:164617 CAPLUS DOCUMENT NUMBER: 132:218003 Nucleic acids encoding human membrane-bound proteins TITLE: and receptors Baker, Kevin; Goddard, Audrey; Gurney, Austin L.; INVENTOR(S): Smith, Victoria; Watanabe, Colin K.; Wood, William I. PATENT ASSIGNEE(S): Genentech, Inc., USA SOURCE: PCT Int. Appl., 773 pp. CODEN: PIXXD2 DOCUMENT TYPE: Patent English LANGUAGE: FAMILY ACC. NUM. COUNT: 143 PATENT INFORMATION: PATENT NO. APPLICATION NO. KIND DATE DATE -----_____ -----_ _ _ _ _____

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ED Entered STN: 13 Mar 2000

AB Membrane-bound proteins and receptor mols. have various industrial applications, including as pharmaceutical and diagnostic agents. Many efforts are focused on the screening of mammalian recombinant DNA libraries to identify the coding sequences for novel receptor or membrane-bound proteins. The present invention is directed to 123 novel polypeptides and to nucleic acid mols. encoding those polypeptides identified in human cDNA libraries by (1) extracellular domain homol. screening, (2) amylase screening, or (3) signal algorithm anal. Also provided herein are vectors and host cells comprising those nucleic acid sequences, chimeric polypeptide mols. comprising the polypeptides of the present invention fused to heterologous polypeptide sequences, antibodies which bind to the polypeptides of the present invention and to methods for producing the polypeptides of the present invention.

IC ICM C12N015-12

ICS C07K014-705; C12N015-62; C07K016-28; C12Q001-68; C12N005-10

CC 3-3 (Biochemical Genetics)

Section cross-reference(s): 6, 13, 63

IT Epitopes

(chimeric proteins containing; nucleic acids encoding human membrane-bound proteins and receptors)

IT Immunoglobulins

RL: BPN (Biosynthetic preparation); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses) (fusion products; nucleic acids encoding human membrane-bound

proteins and receptors)

Fusion proteins (chimeric proteins)
RL: BPN (Biosynthetic preparation); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

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DOCUMENT NUMBER:
                         Cloning and cDNA and deduced amino acid sequences of
TITLE:
                         98 human secreted proteins
                         Komatsoulis, George A.; Rosen, Craig A.; Ruben, Steven
INVENTOR(S):
                         M.; Duan, Roxanne; Moore, Paul A.; Shi, Yanggu;
                         Lafleur, David; Wei, Ying-Fei; Ni, Jian; Florence,
                         Kimberly A.; Young, Paul E.; Brewer, Laurie A.;
                         Soppet, Daniel R.; Endress, Gregory A.; Ebner,
                         Reinhard; Olsen, Henrik S.; Mucenski, Michael
                         Human Genome Sciences, Inc., USA
PATENT ASSIGNEE(S):
                         PCT Int. Appl., 634 pp.
SOURCE:
                         CODEN: PIXXD2
DOCUMENT TYPE:
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LANGUAGE:
                         English
FAMILY ACC. NUM. COUNT:
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PATENT INFORMATION:

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ED
    Entered STN: 11 Feb 2000
AB
    The present invention relates to 98 novel human secreted proteins and
    isolated nucleic acids containing the coding regions of the genes encoding
    such proteins. Tissue distribution, sequence homologies, and preferred
    epitope sites are provided for the secreted proteins, as well as
    chromosomal mapping of some of the genes. Also provided are vectors, host
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- cells, antibodies, and recombinant methods for producing human secreted proteins in bacterial, insect, and mammalian cells. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and treating disorders related to these novel human secreted proteins. High-throughput screening assays are also provided for various putative activities of the secreted proteins.
- IC ICM C12N001-21
 - ICS C12N005-10; C12N015-11; C12N015-12; C12N015-63; A61K038-16; A61K038-17; C07K014-00; C07K014-435; C07K016-00; G01N033-50
- CC 3-3 (Biochemical Genetics)
 - Section cross-reference(s): 6, 13, 63
- IT Immunoglobulins

RL: BPN (Biosynthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(fusion products; cloning and cDNA and deduced amino acid sequences of 98 human secreted proteins)

IT Protein sequences

(of 98 human secreted proteins)

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     ANSWER 45 OF 86
                         2000:68575 CAPLUS
ACCESSION NUMBER:
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DOCUMENT NUMBER:
                          Plant genes encoding cellulose synthase proteins
TITLE:
                         Allen, Stephen M.; Fader, Gary M.; Falco, Saverio
INVENTOR(S):
                          Carl; Kinney, Anthony J.; Lightner, Jonathan E.; Miao,
                         Guo-Hua; Rafalski, J. Antoni; Thorpe, Catherine J.
                          E. I. Du Pont de Nemours & Co., USA
PATENT ASSIGNEE(S):
                          PCT Int. Appl., 94 pp.
SOURCE:
                         CODEN: PIXXD2
DOCUMENT TYPE:
                         Patent
                          English
LANGUAGE:
FAMILY ACC. NUM. COUNT:
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PATENT INFORMATION:
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     PATENT NO.
                         KIND
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01/19/2006 Searched by Alex Waclawiw

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     Entered STN: 28 Jan 2000
ED
AB
     This invention relates to an isolated cDNA fragments encoding cellulose
     synthase proteins from barley, corn, rice, soybean, and wheat. The DNA
     sequences and deduced amino acid sequences are identified by their
     structural homol. to known cellulose synthases from Arabidopsis thaliana
     and Gossypium hirsutum. The invention also relates to the construction of
     a chimeric gene encoding all or a portion of the cellulose synthase
     protein, in sense or antisense orientation, wherein expression of the
     chimeric gene results in production of altered levels of the cellulose
     synthase protein in transformed monocot (corn), dicot (soybean), and
     microbial (Escherichia coli) host cells.
IC
     ICM C12N015-54
         C12N001-21; C12N009-10; C12Q001-48; C12Q001-68
CC
     6-3 (General Biochemistry)
     Section cross-reference(s): 3, 11
IT
     Dicotyledon (Magnoliopsida)
     Escherichia coli
     Monocotyledon (Liliopsida)
        (expression of chimeric genes in; plant genes encoding
        cellulose synthase proteins)
IT
     Protein sequences
        (of plant cellulose synthase proteins)
IT
     Chimeric gene
     RL: BPN (Biosynthetic preparation); BUU (Biological use, unclassified);
     BIOL (Biological study); PREP (Preparation); USES (Uses)
        (plant genes encoding cellulose synthase proteins)
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        (unclaimed protein sequence; plant genes encoding
        cellulose synthase proteins)
     ANSWER 46 OF 86 CAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                        2000:68550 CAPLUS
DOCUMENT NUMBER:
                        132:103779
TITLE:
                        Cloning and cDNA and deduced amino acid sequences of
                        71 human secreted proteins
INVENTOR(S):
                        Ruben, Steven M.; Komatsoulis, George; Duan, Roxanne
```

D.; Rosen, Craig A.; Moore, Paul A.; Shi, Yang-Gu; Lafleur, David W.; Ebner, Reinhard; Olsen, Henrik S.; Brewer, Laurie A.; Florence, Kimberly A.; Young, Paul E.; Mucenski, Michael; Endress, Gregory A.; Soppet, Daniel R.

PATENT ASSIGNEE(S):

Human Genome Sciences, Inc., USA; et al.

SOURCE: PCT Int. Appl., 494 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

51

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

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US	US 6534631					20030318				JS 2	000-	4822	73	20000113					
US	US 2003017500						2003	0123	US 2001-984276					20011029					
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PRIORIT	Y APPL	N. I	NFO	. :						US 1998-92921P					P 19980715				
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									1	WO 1999-US15849					W 19990714 `				
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Entered STN: 28 Jan 2000

- The present invention relates to 71 novel human secreted proteins and AB isolated nucleic acids containing the coding regions of the genes encoding such proteins. Tissue distribution, sequence homologies, and preferred epitope sites are provided for the secreted proteins, as well as chromosomal mapping of some of the genes. Also provided are vectors, host cells, antibodies, and recombinant methods for producing human secreted proteins in bacterial, insect, and mammalian cells. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and treating disorders related to these novel human secreted proteins. High-throughput screening assays are also provided for various putative activities of the secreted proteins.
- ICM C12N015-11

ICS C12N015-00; C12N015-63; C07H021-02; C07H021-04

3-3 (Biochemical Genetics)

Section cross-reference(s): 6, 13, 63

Immunoglobulins

RL: BPN (Biosynthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(fusion products; cloning and cDNA and deduced amino acid sequences of 71 human secreted proteins)

IT Protein sequences

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(of 71 human secreted proteins)
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     255700-69-7
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     RL: PRP (Properties)
        (unclaimed protein sequence; cloning and cDNA and
       deduced amino acid sequences of 71 human secreted proteins)
                              THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS
REFERENCE COUNT:
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L8
    ANSWER 47 OF 86 CAPLUS COPYRIGHT 2006 ACS on STN
                        2000:53914 CAPLUS
ACCESSION NUMBER:
DOCUMENT NUMBER:
                        132:118340
                        Identification of cDNA or its fragment of plant
TITLE:
                        homologs to yeast ADA2 transcription coactivator
                        Cahoon, Rebecca E.; Liu, Zhan-Bin; Odell, Joan T.;
INVENTOR (S):
                        Sakai, Hajime
PATENT ASSIGNEE(S):
                        E. I. Du Pont de Nemours & Co., USA
                        PCT Int. Appl., 42 pp.
SOURCE:
                        CODEN: PIXXD2
DOCUMENT TYPE:
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LANGUAGE:
                        English
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
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PRIORITY APPLN. INFO.:
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                                           WO 1999-US15664
    Entered STN: 23 Jan 2000
ED
    This invention relates to isolated cDNA or its fragment of plant homologs
AB
     to yeast ADA transcription coactivator from corn, rice and soybean. The
     invention also relates to the construction of a chimeric gene encoding all
    or a portion of the transcription coactivator, in sense or antisense
    orientation, wherein expression of the chimeric gene results in production of
    altered levels of the transcription coactivator in a transformed host
    cell.
IC
    ICM C12N015-82
    ICS C12N015-29; C07K014-415; C12N005-10; C12Q001-68
     3-2 (Biochemical Genetics)
CC
    Section cross-reference(s): 6, 11
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Agnes Rooke 10/015,956 ITChimeric gene RL: BPN (Biosynthetic preparation); BUU (Biological use, unclassified); BIOL (Biological study); PREP (Preparation); USES (Uses) (indentification of cDNA or its fragment of plant homologs to yeast ADA2 transcription coactivator) IT Protein sequences cDNA sequences (of gene ADA2 transcription coactivator homolog of corn, rice and soybean; indentification of cDNA or its fragment of plant homologs to yeast ADA2 transcription coactivator) 255856-85-0P 255856-86-1P 255856-87-2P 255856-84-9P IT255856-88-3P 255901-54-3P RL: AGR (Agricultural use); BPN (Biosynthetic preparation); BSU (Biological study, unclassified); PRP (Properties); BIOL (Biological study); PREP (Preparation); USES (Uses) (amino acid sequence; indentification of cDNA or its fragment of plant homologs to yeast ADA2 transcription coactivator) ANSWER 48 OF 86 CAPLUS COPYRIGHT 2006 ACS on STN L82000:23195 CAPLUS ACCESSION NUMBER: 132:90045 DOCUMENT NUMBER: TITLE: Mouse hyaluronate synthase isoforms HAS1, HAS2, and HAS3; chimeric enzymes and substitution mutants of them, and effects on enzymic activities Itano, Naoki; Yoshida, Mamoru; Kimata, Koji INVENTOR(S): Seikagaku Kogyo Co., Ltd., Japan PATENT ASSIGNEE(S): Jpn. Kokai Tokkyo Koho, 30 pp. SOURCE: CODEN: JKXXAF DOCUMENT TYPE: Patent LANGUAGE: Japanese FAMILY ACC. NUM. COUNT: PATENT INFORMATION: PATENT NO. KIND DATE APPLICATION NO. DATE ______ ---------______ JP 1998-193788 JP 2000004886 A2 20000111 19980624 <--PRIORITY APPLN. INFO.: JP 1998-193788 Entered STN: 12 Jan 2000 ED The cDNA sequences encoding 3 isoforms of mouse hyaluronate synthase that AB have different properties have been isolated. A series of chimeric proteins comprised of the N-terminus, the middle region, and the C-terminus of the isoforms; and a series of substitution mutants generated by site-specific mutation of each isoforms are also described. The effects of the mutations on the enzymic activities of HAS are also shown. The DNA sequences encoding the chimeric proteins or substitution mutants are claimed. ICM C12N015-09 IC ICS C12N009-00 7-2 (Enzymes) CCSection cross-reference(s): 13 mouse hyaluronate synthase isoform chimeric protein; ST substitution mutation mouse hyaluronate synthase DNA sequences IT (for chimeric proteins or substitution mutants of mouse hyaluronate synthase isoforms HAS1, HAS2, and HAS3) IT Mouse

(mouse hyaluronate synthase isoforms HAS1, HAS2, and HAS3; chimeric enzymes and substitution mutants of them; and effects

TT

on enzymic activities)

Fusion proteins (chimeric proteins)

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RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
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        chimeric enzymes and substitution mutants of them; and effects
        on enzymic activities)
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        (of chimeric proteins or substitution mutants of mouse
        hyaluronate synthase isoforms HAS1, HAS2, and HAS3)
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     Mutation
        (substitution; mouse hyaluronate synthase isoforms HAS1, HAS2, and
        HAS3; chimeric enzymes and substitution mutants of them; and
        effects on enzymic activities)
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        and HAS3; chimeric enzymes and substitution mutants of them;
        and effects on enzymic activities)
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ТТ
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     BIOL (Biological study); PREP (Preparation)
        (mouse hyaluronate synthase isoforms HAS1, HAS2, and HAS3;
        chimeric enzymes and substitution mutants of them; and effects
        on enzymic activities)
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        HAS1, HAS2, and HAS3; chimeric enzymes and substitution
        mutants of them; and effects on enzymic activities)
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        synthase isoforms HAS1, HAS2, and HAS3; chimeric enzymes and
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     ANSWER 49 OF 86 CAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                         1999:784264 CAPLUS
DOCUMENT NUMBER:
                         132:31785
                         Nucleic acids encoding membrane-bound proteins from
TITLE:
                         human
INVENTOR(S):
                         Baker, Kevin; Chen, Jian; Goddard, Audrey; Gurney,
                         Austin L.; Smith, Victoria; Watanabe, Colin K.; Wood,
                         William I.; Yuan, Jean
PATENT ASSIGNEE(S):
                         Genentech, Inc., USA
                         PCT Int. Appl., 822 pp.
SOURCE:
                         CODEN: PIXXD2
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DOCUMENT TYPE:

Patent

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FAMILY ACC. NUM. COUNT: 143
PATENT INFORMATION:

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Entered STN: 10 Dec 1999 ED

The present invention is directed to 135 polypeptides and to nucleic acid AB mols. encoding those polypeptides. The extracellular domain sequences (including the secretion signal sequence, if any) from about 950 known secreted proteins from the Swiss-Prot public database were used to search EST (expressed sequence tag) databases, and this homol. screen used to assemble consensus DNA sequences relative to other identified EST sequences. Based upon the consensus sequences obtained, oligonucleotides were than synthesized and used to identify by PCR a cDNA library that contained the sequences of interest and for use as probes to isolate clones of full-length coding sequences for the PRO polypeptides. Also provided herein are vectors and host cells comprising those nucleic acid sequences, chimeric polypeptide mols. comprising the polypeptides of the present invention fused to heterologous polypeptide sequences, antibodies which bind to the polypeptides of the present invention and to methods for producing the polypeptides of the present invention. This invention is particularly useful for screening compds. by using PRO polypeptides or binding fragment thereof in any of a variety of drug screening techniques. IC ICM C12N015-12

C07K014-705; C12N015-62; C07K016-28

CC 3-3 (Biochemical Genetics)

Section cross-reference(s): 6, 13

IT Fusion proteins (chimeric proteins)

RL: BPN (Biosynthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses) (nucleic acids encoding membrane-bound proteins from human)

IT Protein sequences

252051-59-5P

(of membrane-bound proteins from human)

252051-62-0P

207624-44-0P, BS106 protein (human fragment) 209859-57-4P 212704-82-0P IT 212838-47-6P 213468-10-1P 213468-15-6P 217795-43-2P, Protein (human clone HP10230) 217795-45-4P, Protein (human clone HP10408) 217795-48-7P, Protein (human clone HP10419) 218948-50-6P 222190-03-6P 222614-92-8P 222618-83-9P 224301-63-7P 220706-50-3P 229477-07-0P 235088-04-7P 243122-35-2P 243646-92-6P, 227792-85-0P Protein (human prostate 371-amino acid) 243976-42-3P 244004-81-7P 249610-95-5P 251314-72-4P, Protein (human clone 586271 244028-85-1P gene Lng107) 251929-74-5P 251929-75-6P 252049-74-4P 252049-78-8P 252049-82-4P 252049-85-7P 252049-87-9P 252049-91-5P 252049-80-2P 252050-03-6P 252050-18-3P 252050-21-8P 252049-94-8P 252050-01-4P 252050-24-1P 252050-26-3P 252050-28-5P 252050-31-0P 252050-35-4P 252050-49-0P 252050-39-8P 252050-43-4P 252050-53-6P 252050-55-8P 252050-75-2P 252050-78-5P 252050-58-1P 252050-61-6P 252050-63-8P 252050-81-0P 252050-83-2P 252050-85-4P 252050-88-7P 252050-92-3P 252051-27-7P 252051-29-9P 252051-00-6P 252051-07-3P 252051-18-6P 252051-43-7P 252051-32-4P 252051-35-7P 252051-38-0P 252051-40-4P 252051-56-2P 252051-45-9P 252051-48-2P 252051-50-6P 252051-54-0P 252052-09-8P

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RL: BOC (Biological occurrence); BPN (Biosynthetic preparation); BSU
(Biological study, unclassified); PRP (Properties); THU (Therapeutic use);
BIOL (Biological study); OCCU (Occurrence); PREP (Preparation); USES
(Uses)
   (amino acid sequence; nucleic acids encoding membrane-bound proteins
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L8 ANSWER 50 OF 86 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1999:736899 CAPLUS

DOCUMENT NUMBER: 132:956

from human)

TITLE: Cloning and cDNA and deduced amino acid sequences of

97 human secreted proteins

INVENTOR(S): Ruben, Steven M.; Florence, Kimberly; Ni, Jian; Rosen,

Craig A.; Carter, Kenneth C.; Moore, Paul A.; Olsen,

Henrik S.; Shi, Yang-Gu; Young, Paul E.; Wei,
Fing-Fei; Brewer, Laurie A.; Soppet, Daniel R.;
Lafleur, David W.; Endress, Gregory A.; Ebner,

Reinhard

PATENT ASSIGNEE(S): Human Genome Sciences, Inc., USA

SOURCE: PCT Int. Appl., 475 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

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ED Entered STN: 19 Nov 1999

The present invention relates to 97 novel human secreted proteins and AB isolated nucleic acids containing the coding regions of the genes encoding such proteins. Tissue distribution, sequence homologies, and preferred epitope sites are provided for the secreted proteins, as well as chromosomal mapping of some of the genes. Also provided are vectors, host cells, antibodies, and recombinant methods for producing human secreted proteins in bacterial, insect, and mammalian cells. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and treating disorders related to these novel human secreted proteins. High-throughput screening assays are also provided for various putative activities of the secreted proteins.

IC ICM C12N015-00

> ICS C12N015-12; C07K014-00; C07K014-435

3-3 (Biochemical Genetics) CC

Section cross-reference(s): 6, 13, 63

IT Immunoglobulins

> RL: BPN (Biosynthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(fusion products; cloning and cDNA and deduced amino acid sequences of 97 human secreted proteins)

IT Protein sequences

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(of 97 human secreted proteins)
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RL: BOC (Biological occurrence); BPN (Biosynthetic preparation); BSU

(Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); OCCU (Occurrence); PREP (Preparation); USES (Uses)

(amino acid sequence; cloning and cDNA and deduced amino acid sequences of 97 human secreted proteins)

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(unclaimed protein sequence; cloning and cDNA and

deduced amino acid sequences of 97 human secreted proteins)

REFERENCE COUNT: 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

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=> fil caplus FILE 'CAPLUS' ENTERED AT 12:17:27 ON 19 JAN 2006 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2006 AMERICAN CHEMICAL SOCIETY (ACS)

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'OBI' IS DEFAULT SEARCH FIELD FOR 'CAPLUS' FILE

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L14 520 SEA FILE=CAPLUS ABB=ON PLU=ON L13 AND L7

L15 143 SEA FILE=CAPLUS ABB=ON PLU=ON L14 AND PY<2001
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L16

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L16 ANSWER 21 OF 125 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2002:461199 CAPLUS

DOCUMENT NUMBER: 137:32059

TITLE: Proteins and genes from Mycobacterium vaccae and

methods for treatment and diagnosis of mycobacterial

infections

INVENTOR(S): Tan, Paul; Visser, Elizabeth; Prestidge, Ross; Watson,

James D.

PATENT ASSIGNEE(S): Genesis Research and Development Corporation Limited,

N.Z.

SOURCE: U.S., 147 pp., Cont.-in-part of U.S. Ser. No. 95,855.

CODEN: USXXAM

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 8

PATENT INFORMATION:

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US 1998-156181

A 19980917

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US 1998-205426
                                                                A 19981204
                                            WO 1998-NZ189
                                                                W 19981223
     Entered STN: 20 Jun 2002
ED
     The present invention provides polypeptides comprising an immunogenic
AB
     portion of a Mycobacterium vaccae protein and DNA mols. encoding such
     polypeptides, together with methods for their use in the diagnosis and
     treatment of mycobacterial infection, including M. tuberculosis and M.
     avium. The invention is further related to compds. that function as
     non-specific immune response amplifiers, and the use of such nonspecific
     immune response amplifiers as adjuvants in vaccination or immunotherapy
     against infectious disease, and in certain treatments for immune disorders
     and cancer. Methods for enhancing the immune response to an antigen
     including administration of M. vaccae culture filtrate, delipidated M.
     vaccae cells, or delipidated and deglycolipidated M. vaccae cells are also
     provided.
     ICM A61K039-04
     ICS
         A61K032-02; G12N001-12
INCL 424248100
     15-2 (Immunochemistry)
     Section cross-reference(s): 3, 6, 10
IT
     Fusion proteins (chimeric proteins)
     RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (proteins and genes from Mycobacterium vaccae and methods for treatment
        and diagnosis of mycobacterial infections)
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     RL: BSU (Biological study, unclassified); PAC (Pharmacological activity);
     PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES
     (Uses)
        (amino acid sequence; proteins and genes from Mycobacterium vaccae and
        methods for treatment and diagnosis of mycobacterial infections)
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     RL: PRP (Properties)
        (unclaimed protein sequence; proteins and genes from Mycobacterium
        vaccae and methods for treatment and diagnosis of mycobacterial
        infections)
REFERENCE COUNT:
                         18
                               THERE ARE 18 CITED REFERENCES AVAILABLE FOR THIS
                               RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
L16 ANSWER 22 OF 125 CAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                         2002:334920 CAPLUS
DOCUMENT NUMBER:
                         136:320425
TITLE:
                         Cloning and characterization of human and murine Rse
                         and HPTK6 receptor protein tyrosine kinases and their
                         antibodies
INVENTOR(S):
                         Godowski, Paul J.; Mark, Melanie R.; Scadden, David T.
PATENT ASSIGNEE(S):
                         Genentech, Inc., USA; New England Deaconess Hospital
```

U.S., 79 pp., Cont. of U.S. Ser. No. 170,558. SOURCE:

CODEN: USXXAM

Patent DOCUMENT TYPE: English LANGUAGE:

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.		DATE
US 5709858	Α	19980120	US 1995-445640		19950522 <
US 6001621	A	19991214	US 1993-170558		19931220 <
CA 2175893	AA	19950601	CA 1994-2175893		19941115 <
US 6087144	Α	20000711	US 1995-447314		19950522 <
US 6096527	Α	20000801	US 1995-445461		19950522 <
US 2002147325	A1	20021010	US 1998-223490		19981230
US 6825324	B2	20041130			
US 2003204072	A1	20031030	US 1999-236939		19990125
US 2004224386	A1	20041111	US 2003-646760		20030825
PRIORITY APPLN. INFO.:			US 1993-157563	В1	19931123
			US 1993-170558	A1	19931220
			US 1998-223490	A1	19981230

ED Entered STN: 06 May 2002

Two protein tyrosine kinase receptors, designated Rse and HPTK6, are AB provided from human and/or murine cell tissues. Rse and HPTK6 were cloned from a cDNA library of a human liver carcinoma cell line (i.e., Hep 3B) using PCR amplification with degenerate oligodeoxyribonucleotide primers designed from sequences encoding conserved amino acids in tyrosine kinases. The murine homolog of Rse was obtained by screening a murine brain cDNA library with a random-primed probe corresponding to nucleotides 1-1163 from human Rse cDNA. Northern blot yielded tissue expression profiles, and the human Rse gene was localized on chromosome 15. Provided herein are nucleic acid sequences encoding Rse and HPTK6 useful as diagnostics and in the recombinant preparation of Rse and HPTK6. Rse and HPTK6 are used in the preparation and purification of antibodies thereto and in diagnostic

assays.

IC ICM A61K039-395

ICS C07K016-28

INCL 424143100

3-3 (Biochemical Genetics)

Section cross-reference(s): 7, 13, 15

IT Human herpesvirus 1

(fusion products with glycoprotein qD of; cloning and characterization of human and murine Rse and HPTK6 receptor protein tyrosine kinases and their antibodies)

ITGlycoproteins

> RL: BPN (Biosynthetic preparation); BUU (Biological use, unclassified); BIOL (Biological study); PREP (Preparation); USES (Uses)

(gD, fusion products with herpes simplex virus 1; cloning and characterization of human and murine Rse and HPTK6 receptor protein tyrosine kinases and their antibodies)

IT 412405-53-9P 412405-55-1P 412405-57-3P 412405-59-5P

412405-61-9P 412405-63-1P

RL: ANT (Analyte); BPN (Biosynthetic preparation); BSU (Biological study, unclassified); DGN (Diagnostic use); PRP (Properties); ANST (Analytical study); BIOL (Biological study); PREP (Preparation); USES (Uses)

(amino acid sequence; cloning and characterization of human and murine Rse and HPTK6 receptor protein tyrosine kinases and their antibodies)

412310-27-1 412310-28-2 412310-29-3 412310-30-6 412310-31-7 IT

RL: PRP (Properties)

(unclaimed sequence; cloning and characterization of human and murine Rse and HPTK6 receptor protein tyrosine kinases and their antibodies) REFERENCE COUNT: 59 THERE ARE 59 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L16 ANSWER 23 OF 125 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2002:153668 CAPLUS

DOCUMENT NUMBER: 136:215389

TITLE:

Vaccine compositions and methods for the prevention and treatment of Mycobacterium tuberculosis infection Reed, Steven G.; Skeiky, Yasir A. W.; Dillon, Davin C. INVENTOR(S):

PATENT ASSIGNEE(S): Corixa Corporation, USA

SOURCE: U.S., 156 pp., Cont.-in-part of U.S. Ser. No. 25,197,

> abandoned. CODEN: USXXAM

DOCUMENT TYPE: Patent English LANGUAGE:

FAMILY ACC. NUM. COUNT: 13

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PR:

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US 1998-223040 A 19981230
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WO 1999-US7717 W 19990407

ED Entered STN: 28 Feb 2002

AB Compns. and methods for treatment and vaccination against tuberculosis are disclosed. In one aspect the compns. provided include at least two polypeptides that contain an immunogenic portion of a M. tuberculosis antigen or at least two DNA mols. encoding such polypeptides. In a second aspect, the compns. provided include a fusion protein comprising at least two polypeptides that contain an immunogenic portion of a M. tuberculosis antigen. Such compns. may be formulated into vaccines and/or pharmaceutical compns. for immunization against M. tuberculosis infection, or may be used for the diagnosis of tuberculosis. The vaccine compns. may also comprises immune adjuvant such as 3D-MPL or QS21 and immunostimulatory cytokine or chemokine, and is formulated in oil-in-water emulsion.

IC ICM A61K003-04

ICS A61K039-40; C12Q001-68; G01N033-53; C12N001-12

INCL 424248100

CC 15-2 (Immunochemistry)

Section cross-reference(s): 3, 63

IT Fusion proteins (chimeric proteins)

RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified); DGN (Diagnostic use); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(Mycobacterium tuberculosis antigens for diagnosis and treatment of tuberculosis)

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RL: ANT (Analyte); BPN (Biosynthetic preparation); BSU (Biological study, unclassified); DGN (Diagnostic use); PRP (Properties); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); PREP (Preparation); USES (Uses)

(amino acid sequence; mycobacterium tuberculosis antigens for diagnosis and treatment of tuberculosis)

RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

and treatment of tuberculosis)
REFERENCE COUNT: 10 THERE ARE 10 CITED REFERENCES AVAILABLE FOR THIS

L16 ANSWER 24 OF 125 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2002:90102 CAPLUS

DOCUMENT NUMBER: 136:146183

TITLE: Nucleic acid and protein compositions and methods for the diagnosis and treatment of disorders involving

angiogenesis

INVENTOR(S):

Baker, Kevin P.; Ferrara, Napoleone; Gerber,
Hanspeter; Gerritsen, Mary E.; Goddard, Audrey;
Godowski, Paul J.; Gurney, Austin L.; Hillan, Kenneth
J.; Marsters, Scot A.; Pan, James; Paoni, Nicholas F.;
Stephan, Jean-Philippe F.; Watanabe, Colin K.;
Williams, P. Mickey; Wood, William I.; Ye, Weilan

PATENT ASSIGNEE(S): SOURCE:

Genentech, Inc., USA PCT Int. Appl., 567 pp.

CODEN: PIXXD2

DOCUMENT TYPE: LANGUAGE: Patent English

FAMILY ACC. NUM. COUNT: 143

PATENT INFORMATION:

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WO 1999-US21090
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WO 1999-US21547
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ED Entered STN: 01 Feb 2002

Nucleic acid and protein compns. and methods are disclosed for stimulating AB or inhibiting angiogenesis and/or cardiovascularization in mammals, including humans. Thus, 187 cDNAs and their encoded protein sequences isolated from human cDNA libraries are identified by extracellular domain homol. screening, amylase screening, and signal algorithm anal. The pharmaceutical compns. are based on polypeptides or antagonists thereto that have been identified for one or more of these uses. Disorders that can be diagnosed, prevented, or treated by the compns. herein include trauma such as wounds, various cancers, and disorders of the vessels including atherosclerosis and cardiac hypertrophy. Also provided herein are vectors and host cells comprising those nucleic acid sequences, chimeric polypeptide mols. comprising the polypeptides of the present invention fused to heterologous polypeptide sequences, antibodies which bind to the polypeptides of the present invention, and to methods for producing the polypeptides of the present invention.

IC C07K014-475

CC 3-3 (Biochemical Genetics)

Section cross-reference(s): 6, 9, 13, 63

IT Epitopes

(chimeric proteins containing; nucleic acid and protein compns. and methods for the diagnosis and treatment of disorders involving angiogenesis)

IT Antibodies and Immunoglobulins

RL: BPN (Biosynthetic preparation); DGN (Diagnostic use); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(fragments, chimeric proteins containing Fc region; nucleic acid and protein compns. and methods for the diagnosis and treatment of disorders involving angiogenesis)

IT Fusion proteins (chimeric proteins)

RL: BPN (Biosynthetic preparation); DGN (Diagnostic use); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(nucleic acid and protein compns. and methods for the diagnosis and treatment of disorders involving angiogenesis)

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IT

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(Biological study); PREP (Preparation); USES (Uses)
   (amino acid sequence; nucleic acid and protein compns. and methods for
   the diagnosis and treatment of disorders involving angiogenesis)
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L16 ANSWER 25 OF 125 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER:

2002:72710 CAPLUS

DOCUMENT NUMBER:

136:149852

TITLE:

Fusion proteins of Mycobacterium

tuberculosis antigens and their uses

INVENTOR (S):

Reed, Steven G.; Skeiky, Yasir A.; Dillon, Davin C.;

Alderson, Mark; Campos-Neto, Antonio

PATENT ASSIGNEE(S):

USA

SOURCE:

U.S. Pat. Appl. Publ., 62 pp., Cont.-in-part of U.S.

Ser. No. 223,040.

CODEN: USXXCO

DOCUMENT TYPE:

Patent English

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2002009459	A1	20020124	US 1999-287849	19990407
US 6627198	B2	20030930		
US 6290969	B1	20010918	US 1997-818112	19970313
US 6350456	B1	20020226	US 1998-56556	19980407
US 6544522	B1	20030408	US 1998-223040	19981230
ZA 9901303	Α	20000315	ZA 1999-1303	19990218 <
TW 223001	B1	20041101	TW 1999-88105532	19990426
AU 765833	B2	20031002	AU 2000-71762	20001122
US 2003147911	A1	20030807	US 2003-359460	20030205
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US 2003-359459
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PRIORITY APPLN. INFO.:
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                                            US 1999-287849
                                                                B1 19990407
    Entered STN: 27 Jan 2002
ED
    The present invention relates to fusion proteins containing at least two
AB
    Mycobacterium tuberculosis antigens. In particular, it relates to
    bi-fusion proteins which contain two individual M. tuberculosis antigens,
     tri-fusion proteins which contain three M. tuberculosis antigens,
     tetra-fusion proteins which contain four M. tuberculosis antigens, and
     penta-fusion proteins which contain five M. tuberculosis antigens, and
     methods for their use in the diagnosis, treatment and prevention of
     tuberculosis infection.
     ICM A61K039-02
TC
INCL 424190100
     15-2 (Immunochemistry)
     Section cross-reference(s): 3
    Mycobacterium tuberculosis antigen fusion protein vaccine
ST
     Cell proliferation
IT
        (T cell; fusion proteins of Mycobacterium tuberculosis
        antigens and their uses)
TТ
     Immunostimulants
        (adjuvants; fusion proteins of Mycobacterium tuberculosis
        antigens and their uses)
     Immunity
IT
        (cell-mediated; fusion proteins of Mycobacterium tuberculosis
        antigens and their uses)
IT
    B cell (lymphocyte)
    DNA sequences
    Molecular cloning
    Mycobacterium tuberculosis
    Nucleic acid hybridization
     Protein sequences
    T cell (lymphocyte)
     Tuberculosis
     Vaccines
        (fusion proteins of Mycobacterium tuberculosis antigens and
        their uses)
IT
    Antigens
       Fusion proteins (chimeric proteins)
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     DGN (Diagnostic use); PRP (Properties); THU (Therapeutic use); BIOL
     (Biological study); PREP (Preparation); USES (Uses)
        (fusion proteins of Mycobacterium tuberculosis antigens and
        their uses)
     Cytokines
IT
     Interleukin 4
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (fusion proteins of Mycobacterium tuberculosis antigens and
        their uses)
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Antibodies and Immunoglobulins
IT
    Polynucleotides
    RL: BSU (Biological study, unclassified); DGN (Diagnostic use); THU
     (Therapeutic use); BIOL (Biological study); USES (Uses)
        (fusion proteins of Mycobacterium tuberculosis antigens and
       their uses)
IT
    Immunity
        (humoral; fusion proteins of Mycobacterium tuberculosis
       antigens and their uses)
    Animal
ΤT
        (non-human; fusion proteins of Mycobacterium tuberculosis
       antigens and their uses)
    T cell (lymphocyte)
IT
        (proliferation; fusion proteins of Mycobacterium tuberculosis
       antigens and their uses)
IT
    DNA
    RL: BSU (Biological study, unclassified); DGN (Diagnostic use); THU
     (Therapeutic use); BIOL (Biological study); USES (Uses)
        (recombinant; fusion proteins of Mycobacterium tuberculosis
       antigens and their uses)
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        (amino acid sequence; fusion proteins of Mycobacterium
        tuberculosis antigens and their uses)
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     (Biological study); PREP (Preparation); USES (Uses)
        (nucleotide sequence; fusion proteins of Mycobacterium
        tuberculosis antigens and their uses)
L16 ANSWER 26 OF 125 CAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                        2002:41632 CAPLUS
DOCUMENT NUMBER:
                        136:117361
                        Stress proteins as immunomodulators and in vaccines as
TITLE:
                        fusion proteins with antigens
                        Young, Richard A.
INVENTOR(S):
                        Whitehead Institute for Biomedical Research, USA
PATENT ASSIGNEE(S):
                        U.S., 29 pp., Cont.-in-part of WO9429459.
SOURCE:
                        CODEN: USXXAM
DOCUMENT TYPE:
                        Patent
                        English
LANGUAGE:
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
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                                         APPLICATION NO.
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     WO 8912455
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                                            WO 1994-US6362
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     EP 1221488
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                                20020710
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        R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE
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     US 2005163792
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PRIORITY APPLN. INFO.:
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                                                                B2 19880615
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                                            WO 1989-US2619
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                                                                B1 20020114
ED
     Entered STN: 16 Jan 2002
AB
     The present invention relates to stress proteins and methods of modulating
     an individual's immune response. In particular, it relates to the use of
     such stress proteins in immune therapy and prophylaxis, which results in
     an induction or enhancement of an individual's immune response and as an
     immunotherapeutic agent which results in a decrease of an individual's
     immune response to his or her own cells. The present invention also
     relates to compns. comprising a stress protein joined to another
     component, such as a fusion protein in which a stress protein is fused to
     an antigen. Further, the present invention relates to a method of
     generating antibodies to a substance using a conjugate comprised of a
     stress protein joined to the substance.
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IC ICM C12P021-04

ICS A61K039-21; A61K039-04; C12N015-00

INCL 435069700

CC 15-2 (Immunochemistry)

Section cross-reference(s): 9, 63

- ST stress protein chimera immunomodulator vaccine
- IT Proteins

RL: BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(65kDa; stress proteins as immunomodulators and in vaccines as **fusion** proteins with antigens)

IT Proteins

RL: BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(71kDa; stress proteins as immunomodulators and in vaccines as **fusion** proteins with antigens)

IT Molecular chaperones

RL: BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(DnaJ; stress proteins as immunomodulators and in vaccines as **fusion** proteins with antigens)

IT Molecular chaperones

RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(DnaK; stress proteins as immunomodulators and in vaccines as **fusion** proteins with antigens)

IT Molecular chaperones

RL: BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (GroEL; stress proteins as immunomodulators and in vaccines as fusion proteins with antigens) IT Molecular chaperones RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (GroES; stress proteins as immunomodulators and in vaccines as fusion proteins with antigens) Heat-shock proteins TΤ RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (HSP 60; stress proteins as immunomodulators and in vaccines as fusion proteins with antigens) IT Heat-shock proteins RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (HSP 65; stress proteins as immunomodulators and in vaccines as fusion proteins with antigens) IT Heat-shock proteins RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (HSP 70; stress proteins as immunomodulators and in vaccines as fusion proteins with antigens) IT Heat-shock proteins RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (HSP 90; stress proteins as immunomodulators and in vaccines as fusion proteins with antigens) IT Proteins RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (P1; stress proteins as immunomodulators and in vaccines as fusion proteins with antigens) IT Immunostimulants (adjuvants; stress proteins as immunomodulators and in vaccines as fusion proteins with antigens) TΤ Antigens RL: BSU (Biological study, unclassified); BIOL (Biological study) (autoantigens; stress proteins as immunomodulators and in vaccines as fusion proteins with antigens) Antibodies and Immunoglobulins IT RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (monoclonal; stress proteins as immunomodulators and in vaccines as fusion proteins with antigens) IT gag proteins RL: BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (p24gag; stress proteins as immunomodulators and in vaccines as fusion proteins with antigens) IT Polyproteins RL: BSU (Biological study, unclassified); BIOL (Biological study) (pol; stress proteins as immunomodulators and in vaccines as fusion proteins with antigens) TT Autoimmune disease Chromatography Escherichia coli Eubacteria

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Fungi
     Human
     Human immunodeficiency virus
     Immune tolerance
     Immunotherapy
     Mammalia
     Microorganism
     Mycobacterium
     Mycobacterium bovis
     Mycobacterium leprae
     Mycobacterium tuberculosis
     Parasite
     Pathogen
     Protein sequences
     Rheumatoid arthritis
     Vaccines
     Vertebrata
     Virus
        (stress proteins as immunomodulators and in vaccines as fusion
        proteins with antigens)
     Antibodies and Immunoglobulins
IT
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES
     (Uses)
        (stress proteins as immunomodulators and in vaccines as fusion
        proteins with antigens)
IT
     gag proteins
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (stress proteins as immunomodulators and in vaccines as fusion
        proteins with antigens)
IT
     Antiqens
       Fusion proteins (chimeric proteins)
     Gene, microbial
     Heat-shock proteins
     Toxins
     Tumor antigens
     RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (stress proteins as immunomodulators and in vaccines as fusion
        proteins with antigens)
IT
     Proteins
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     (Biological study); USES (Uses)
        (stress-induced; stress proteins as immunomodulators and in vaccines as
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     389998-15-6, Protein P1 (human) 389998-16-7, Chaperonin
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        vaccines as fusion proteins with antigens)
REFERENCE COUNT:
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                               THERE ARE 46 CITED REFERENCES AVAILABLE FOR THIS
                               RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
L16 ANSWER 27 OF 125 CAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                         2001:895574 CAPLUS
DOCUMENT NUMBER:
                         136:52707
TITLE:
                         Methods for the treatment of immunologically-mediated
                         skin disorders
INVENTOR(S):
                         Watson, James D.; Tan, Paul L. J.; Prestidge, Ross
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Agnes Rooke 10/015,956 Genesis Research & Development Corp. Ltd., N. Z. PATENT ASSIGNEE(S): U.S., 116 pp., Cont.-in-part of U.S. 5,968,524. SOURCE: CODEN: USXXAM DOCUMENT TYPE: Patent English LANGUAGE: FAMILY ACC. NUM. COUNT: PATENT INFORMATION:

PATENT NO. KIND DATE APPLICATION NO. DATE

US 6328978 B1 20011211 US 1999-324542 19990602
US 5968524 A 19991019 US 1997-997080 19971223 <-IN 188709 A 20021026 IN 2000-CA231 20000419
US 2003007976 A1 20030109 US 2001-880505 20010613 US 2001-880505 20010613 US 1997-997080 A2 19971223 IN 1998-CA242 A 1900000 PRIORITY APPLN. INFO.: IN 1998-CA242 A 19980216 US 1999-324542 A2 19990602

Entered STN: 12 Dec 2001 ED

Methods for the treatment of skin disorders, including psoriasis, atopic AB dermatitis, allergic contact dermatitis, alopecia areata and skin cancers are provided, such methods comprising administering a composition having antigenic and/or adjuvant properties. Compns. which may be usefully employed in the inventive methods include inactivated M. vaccae cells, delipidated and deglycolipidated M. vaccae cells, M. vaccae culture filtrate and compds. present in or derived therefrom, together with combinations of such compns.

ICM A61K045-00

ICS A61K039-04; A61K039-02; A61K039-38; A61K038-00

INCL 424282100

15-2 (Immunochemistry)

Section cross-reference(s): 3, 63

Fusion proteins (chimeric proteins)

RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified); BUU (Biological use, unclassified); BIOL (Biological study); PREP (Preparation); USES (Uses)

(inactivated or delipidated and deglycolipidated Mycobacterium vaccae or antigens for treatment of immunol.-mediated skin disorders)

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RL: BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(amino acid sequence; inactivated or delipidated and deglycolipidated Mycobacterium vaccae or antigens for treatment of immunol.-mediated skin disorders)

IT380693-94-7 380693-95-8 380693-96-9 380693-97-0 380694-01-9 380694-02-0 380694-03-1 380694-04-2 380694-05-3 380694-17-7 **380694-19-9** 380694-26-8 380694-44-0 380694-45-1 380694-48-4 380694-49-5 380694-52-0

RL: PRP (Properties)

(unclaimed protein sequence; methods for the treatment of immunol.-mediated skin disorders)

THERE ARE 24 CITED REFERENCES AVAILABLE FOR THIS REFERENCE COUNT: 24

RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

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L16 ANSWER 28 OF 125 CAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                        2001:771048 CAPLUS
                        135:328149
DOCUMENT NUMBER:
TITLE:
                        Protein and cDNA sequences of human, mouse, and Danio
                        rerio retinoic acid-metabolizing protein, and uses
                        thereof
INVENTOR (S):
                        Petkovich, P. Martin; White, Jay A.; Beckett, Barbara
                        R.; Jones, Glenville
PATENT ASSIGNEE(S):
                        Queen's University at Kingston, Can.
                        U.S., 75 pp., Cont.-in-part of Appl. No.
SOURCE:
                        PCT/CA97/00440.
                        CODEN: USXXAM
DOCUMENT TYPE:
                        Patent
                        English
LANGUAGE:
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
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                                                             A3 20000925
     Entered STN: 24 Oct 2001
ED
    This invention provides protein and cDNA sequences of human, mouse, and
AB
     zebrafish (Danio rerio) retinoic acid-metabolizing protein. The protein
     is shown to have the ability to hydroxylate retinoic acid (RA) at the 4
     position of the \beta-ionone ring. The protein of the invention belongs
     to the family of cytochrome P450s, and its production in epithelial cells is
     induced by treatment with RA. The invention also relates to the use of
     the provided proteins and cDNAs, particularly in drug screening assays.
     The examples disclose studies of the effects of 4-Hydroxyphenylretinamide,
     RA, ketoconazole, and Am580 upon the expression of the protein.
IC
     ICM C12P021-04
     ICS C12N009-02; C12N005-00; C07H021-04
INCL 435069700
    3-3 (Biochemical Genetics)
    Section cross-reference(s): 6, 13, 63
IT
    Fusion proteins (chimeric proteins)
    RL: BPN (Biosynthetic preparation); BIOL (Biological study); PREP
     (Preparation)
        (comprising retinoic acid-metabolizing proteins; protein and cDNA
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sequences of human, mouse, and Danio rerio retinoic acid-metabolizing

protein, and uses thereof) 184722-39-2P 346004-25-9P 368955-04-8P TT RL: BAC (Biological activity or effector, except adverse); BOC (Biological occurrence); BPN (Biosynthetic preparation); BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); OCCU (Occurrence); PREP (Preparation); USES (Uses) (amino acid sequence; protein and cDNA sequences of human, mouse, and Danio rerio retinoic acid-metabolizing protein, and uses thereof) REFERENCE COUNT: THERE ARE 64 CITED REFERENCES AVAILABLE FOR THIS 64 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT L16 ANSWER 29 OF 125 CAPLUS COPYRIGHT 2006 ACS on STN 2001:693506 CAPLUS ACCESSION NUMBER: DOCUMENT NUMBER: 135:268240 Secreted and transmembrane polypeptides and human TITLE: nucleic acids encoding them that are overexpressed in cancerous tissues Baker, Kevin P.; Chen, Jian; Desnoyers, Luc; Goddard, INVENTOR(S): Audrey; Godowski, Paul J.; Gurney, Austin L.; Pan, James; Smith, Victoria; Watanabe, Colin K.; Wood, William I.; Zhang, Zemin Genentech, Inc., USA PATENT ASSIGNEE(S): PCT Int. Appl., 774 pp. SOURCE: CODEN: PIXXD2 Patent DOCUMENT TYPE: English LANGUAGE: FAMILY ACC. NUM. COUNT: 143 PATENT INFORMATION:

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                   A 20010309
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ED Entered STN: 21 Sep 2001

The present invention is directed to novel polypeptides and to nucleic acid mols. encoding those polypeptides. Thus, 305 cDNAs encoding human secreted or transmembrane proteins were identified by extracellular domain homol. screening, amylase screening, and signal algorithm anal. These transcripts for these proteins are overexpressed in various cancerous tissues, including adrenal, lung, colon, breast, prostate, rectal, cervical, and liver tumors. Certain of the proteins stimulate release of tumor necrosis factor- α from human blood, and also stimulate proliferation or differentiation of chondrocytes. Also provided herein are vectors and host cells comprising those nucleic acid sequences, chimeric polypeptide mols. comprising the polypeptides of the present invention fused to heterologous polypeptide sequences, antibodies which bind to the polypeptides of the present invention and to methods for producing the polypeptides of the present invention.

IC ICM C12N015-12

ICS C12N015-62; C07K014-47; C07K014-705; C07K016-18; G01N033-53; C12O001-68

CC 3-3 (Biochemical Genetics)

Section cross-reference(s): 6, 13, 14

IT Antibodies

RL: BPN (Biosynthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses) (chimeric; secreted and transmembrane polypeptides and human

(chimeric; secreted and transmembrane polypeptides and human nucleic acids encoding them that are overexpressed in cancerous tissues)

IT Fusion proteins (chimeric proteins)

RL: BPN (Biosynthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(secreted and transmembrane polypeptides and human nucleic acids encoding them that are overexpressed in cancerous tissues)

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RL: ANT (Analyte); BOC (Biological occurrence); BSU (Biological study,
unclassified); PRP (Properties); THU (Therapeutic use); ANST (Analytical
study); BIOL (Biological study); OCCU (Occurrence); USES (Uses)
   (amino acid sequence; secreted and transmembrane polypeptides and human
   nucleic acids encoding them that are overexpressed in cancerous
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RL: ANT (Analyte); BOC (Biological occurrence); BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); OCCU (Occurrence); USES (Uses) (amino acid sequence; secreted and transmembrane polypeptides and human nucleic acids encoding them that are overexpressed in cancerous

L16 ANSWER 30 OF 125 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2001:611697 CAPLUS

DOCUMENT NUMBER: 135:192174

TITLE: Sequence, structure and interaction with TRF1 of human

tankyrase and potential use of tankyrase in drug

screening

INVENTOR(S): De Lange, Titia; Smith, Susan
PATENT ASSIGNEE(S): Rockefeller University, USA

SOURCE: U.S., 65 pp., Cont.-in-part of U.S. Ser. No. 135,223.

CODEN: USXXAM

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2

tissues)

PATENT INFORMATION:

PATENT NO.						APPLICATION NO.					DATE						
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	6277																
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ED Entered STN: 23 Aug 2001

AB The present invention discloses cDNA and amino acid sequences of human tankyrase that binds to the N-terminal acidic domain of telomeric repeat binding factor 1 (TRF1). A full-length human tankyrase is 142-kDa protein and contains an ankyrin-specific repeat consensus domain, a sterile alpha motif (SAM), and a poly (ADP-ribose) polymerase-related domain. Amino acid and encoding cDNA sequence of two truncated forms of human tankyrase are also disclosed. Methods of screening drugs using tankyrase are included.

IC ICM C12N009-10

INCL 435193000

CC 7-5 (Enzymes)

Section cross-reference(s): 1, 3, 13

IT Fusion proteins (chimeric proteins)

RL: BPN (Biosynthetic preparation); BUU (Biological use, unclassified); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(tankyrase contg; sequence, structure and interaction with TRF1 of
human tankyrase and potential use of tankyrase in drug screening)
252647-70-4P 252647-87-3P 355174-12-8DP, substitution mutants are
claimed 355174-13-9DP, substitution mutants are claimed
355174-14-0DP, substitution mutants are claimed 355174-15-1DP,
substitution mutants are claimed 355174-16-2DP, substitution mutants are
claimed 355174-17-3DP, substitution mutants are claimed 355174-18-4DP,
substitution mutants are claimed 355387-93-8DP, substitution mutants are
claimed

RL: BPN (Biosynthetic preparation); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(amino acid sequence; sequence, structure and interaction with TRF1 of human tankyrase and potential use of tankyrase in drug screening)

9055-67-8P, Tankyrase 219943-19-8DP, subfragments are claimed RL: BAC (Biological activity or effector, except adverse); BOC (Biological occurrence); BPN (Biosynthetic preparation); BPR (Biological process); BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); OCCU (Occurrence); PREP (Preparation); PROC (Process); USES (Uses)

(sequence, structure and interaction with TRF1 of human tankyrase and potential use of tankyrase in drug screening)

REFERENCE COUNT: 36 THERE ARE 36 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L16 ANSWER 31 OF 125 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2001:482174 CAPLUS

DOCUMENT NUMBER: 135:88006

TITLE: DNA encoding glycogenin involved in starch

biosynthesis

INVENTOR(S): Lightner, Jonathan Edward; Everard, John D. PATENT ASSIGNEE(S): E. I. du Pont de Nemours & Company, USA

SOURCE: U.S., 33 pp., Cont.-in-part of U.S. Ser. No. 852,615,

abandoned.
CODEN: USXXAM

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 3

PATENT INFORMATION:

ΙT

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE		
US 6255114	B1	20010703	US 1998-73297	19980506		
ZA 9803846	Α	19991108	ZA 1998-3846	19980507 <		
US 2002001843	A1	20020103	US 2001-829482	20010410		
US 2003145353	A1	20030731	US 2003-336587	20030102		
PRIORITY APPLN. INFO.:			US 1997-852615 B2	19970507		
			US 1998-73297 A3	19980506		
			US 2001-829482 A2	20010410		

ED Entered STN: 05 Jul 2001

AB This invention relates to isolated nucleic acid fragments encoding all or a substantial portion of a plant glycogenin or water stress protein from corn, rice and wheat. The invention also relates to the construction of chimeric genes encoding all or a portion of a plant glycogenin or water stress protein, in sense or antisense orientation. Expression of the chimeric gene results in production of altered levels of a plant glycogenin or

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water stress protein in a transformed host cell.
IC
     ICM C12N015-29
     ICS C12N005-04; C12N015-74; C12N015-82; C12P019-04
INCL 435468000
     3-3 (Biochemical Genetics)
     Section cross-reference(s): 7, 11
IT
     Chimeric gene
     EST (expressed sequence tag)
     RL: AGR (Agricultural use); BIOL (Biological study); USES (Uses)
        (DNA encoding glycogenin involved in starch biosynthesis)
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IT
     216008-82-1
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     216008-91-2
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                                                             216009-16-4
     216101-28-9
     RL: AGR (Agricultural use); ANT (Analyte); PRP (Properties); ANST
     (Analytical study); BIOL (Biological study); USES (Uses)
        (amino acid sequence; DNA encoding glycogenin involved in starch
        biosynthesis)
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INVENTOR(S):
                         Audrey; Godowski, Paul J.; Grimaldi, Christopher J.;
                         Gurney, Austin L.; Li, Hanzhong; Hillan, Kenneth J.;
                         Tumas, Daniel; Van, Lookeren Menno; Vandlen, Richard
                         L.; Watanabe, Colin K.; Williams, P. Mickey; Wood,
                         William I.; Yansura, Daniel G.
                         Genentech, Inc., USA
PATENT ASSIGNEE(S):
                         PCT Int. Appl., 198 pp.
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ED

Entered STN: 29 Jun 2001 Proteins that are sequence homologs of interleukin 17 and that therefore AΒ may be of use in the treatment of diseases associated with immune dysfunction are described. Cloning and expression vectors and host cells, fusion

proteins, antibodies to the proteins, and methods of manufacturing them are described. The genes for these proteins are expressed in a wide array of tissues. The proteins do not bind known interleukin 17 receptors but one of the proteins was identified as a receptor for two of the others. The proteins show interleukin 17-like activities, such as activation of NF- κ B.

IC ICM C12N015-12

ICS C07K014-715; C07K014-54; C12N015-62; C07K016-24; C07K016-28;
 A61K031-70; A61K038-17; A61K038-20; A61K039-395; G01N033-566;
 G01N033-53

CC 15-5 (Immunochemistry)

Section cross-reference(s): 1, 3

IT Fusion proteins (chimeric proteins)

RL: BPN (Biosynthetic preparation); THU (Therapeutic use); BIOL

(Biological study); PREP (Preparation); USES (Uses)

(homologs of interleukin 17 identified by sequence similarity and their use in treatment of immune dysfunction)

IT 187759-17-7 215663-51-7 251100-02-4, Interleukin 21 (human)

321202-67-9 329335-25-3 329800-21-7 347432-44-4

347432-47-7 347432-49-9

RL: BPR (Biological process); BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PROC (Process); USES (Uses)

(amino acid sequence; homologs of interleukin 17 identified by sequence similarity and their use in treatment of immune dysfunction)

L16 ANSWER 33 OF 125 CAPLUS COPYRIGHT 2006 ACS on STN

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INVENTOR(S): Labigne, Agnes; Suerbaum,
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PATENT ASSIGNEE(S): Institut Pasteur, Fr.

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                                             WO 1996-EP1834
     Entered STN: 21 Jun 2001
ED
     There is provided an immunogenic composition capable of inducing protective
AB
     antibodies against Helicobacter infection characterized in that it
     comprises: i) at least one sub-unit of a urease structural polypeptide
     from Helicobacter pylori, or a fragment thereof, said fragment being
     recognized by antibodies reacting with Helicobacter felis urease, and/or
     at least one sub-unit of a urease structural polypeptide from Helicobacter
     felis, or a fragment thereof, said fragment being recognized by antibodies
     reacting with Helicobacter pylori urease; ii) and/or, a heat shock protein
     (Hsp), or chaperonin, from Helicobacter, or a fragment of said protein.
     The preparation, by recombinant means, of such immunogenic compns. is also
     provided.
IC
     A61K039-00
INCL 424192100
     15-2 (Immunochemistry)
     Section cross-reference(s): 3, 9, 10
IT
     Proteins, specific or class
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES
     (Uses)
        (MBP (maltose-binding protein), chimeric; immunogenic compns.
        comprising Helicobacter heat shock protein or chaperonin for vaccine
        against Helicobacter infection)
IT
     Antigens
       Fusion proteins (chimeric proteins)
     Heat-shock proteins
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
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     (Preparation); USES (Uses)
        (immunogenic compns. comprising Helicobacter heat shock protein or
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     162243-40-5, Urease (Helicobacter felis gene ureA)
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     115681-99-7, Protein (Escherichia coli gene groES) 117537-95-8 127314-53-8, Urease (Proteus mirabilis clone pMID1003 \gamma-subunit
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                               142193-37-1, Protein (Helicobacter pylori clone
     protein moiety reduced)
     pILL753 gene ureI reduced)
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     perfringens clone pCPH-2 gene groES) 162243-42-7, Urease (Helicobacter
     felis gene ureI) 244773-35-1
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     345920-31-2
     345920-35-6
     RL: PRP (Properties)
        (unclaimed protein sequence; immunogenic compns. against Helicobacter
        infection, polypeptides for use in the compns. and nucleic acid
        sequences encoding said polypeptides)
                               THERE ARE 62 CITED REFERENCES AVAILABLE FOR THIS
REFERENCE COUNT:
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L16 ANSWER 34 OF 125 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2001:417147 CAPLUS

DOCUMENT NUMBER: 135:29838

TITLE: Secreted and transmembrane proteins identified by

sequence comparison and cDNAs encoding them and their

uses

INVENTOR(S): Baker, Kevin; Beresini, Maureen; Deforge, Laura;

Desnoyers, Luc; Filvaroff, Ellen; Gao, Wei Qiang; Gerritsen, Amry E.; Goddard, Audrey; Godowski, Paul J.; Gurney, Austin L.; Gherwood, Steven; Smith, Victoria; Stewart, Timothy A.; Tumas, Daniel;

Watanabe, Colin K.; Wood, William I.; Zhang, Zemin

PATENT ASSIGNEE(S): Genentech, Inc., USA SOURCE: PCT Int. Appl., 813 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 143

PATENT INFORMATION:

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ED Entered STN: 08 Jun 2001

AB The present invention is directed to novel polypeptides and to nucleic acid mols. encoding those polypeptides. Also provided herein are vectors and host cells comprising those nucleic acid sequences, chimeric polypeptide mols. comprising the polypeptides of the present invention fused to heterologous polypeptide sequences, antibodies which bind to the polypeptides of the present invention and to methods for producing the polypeptides of the present invention. The proteins show overexpression in cancer and may of diagnostic use. Certain of the proteins were found to form complexes with one another.

IC C12N015-12; C07K014-47

CC 3-2 (Biochemical Genetics)

Section cross-reference(s): 6

IT Immunoglobulins

RL: BPN (Biosynthetic preparation); BIOL (Biological study); PREP (Preparation)

(fusion products, with secreted and transmembrane proteins of human; secreted and transmembrane proteins identified by sequence comparison and cDNAs encoding them and their uses)

IT Immunoglobulins

RL: BPN (Biosynthetic preparation); BIOL (Biological study); PREP (Preparation)

(heavy chains, constant region C-terminal fragment, fusion products with secreted and transmembrane proteins of human; secreted

and transmembrane proteins identified by sequence comparison and cDNAs encoding them and their uses)

IT Fusion proteins (chimeric proteins)

RL: BPN (Biosynthetic preparation); BIOL (Biological study); PREP (Preparation)

(of secreted and transmembrane proteins of human; secreted and transmembrane proteins identified by sequence comparison and cDNAs encoding them and their uses)

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RL: BSU (Biological study, unclassified); PRP (Properties); THU
(Therapeutic use); BIOL (Biological study); USES (Uses)
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L16 ANSWER 35 OF 125 CAPLUS COPYRIGHT 2006 ACS on STN
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ACCESSION NUMBER:
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                         135:4471
TITLE:
                         Antibodies
                         Kingsman, Alan; Kingsman, Susan Mary; Bebbington,
INVENTOR(S):
                         Christopher Robert; Carroll, Miles William; Ellard,
                         Fiona Margaret; Myers, Kevin Alan
PATENT ASSIGNEE(S):
                         Oxford Biomedica (UK) Limited, UK
SOURCE:
                         PCT Int. Appl., 117 pp.
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DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT: 5

PATENT INFORMATION:

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Agnes Rooke 10/015,956
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                                       WO 2001-GB5004
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                                       US 2002-60585
                                                           A2 20020129
Entered STN: 27 May 2001
The use of an ScFv Ab (ScFv Ab) capable of recognizing a disease associated
mol. (DAM) in the manufacture of a medicament for the prevention and/or
treatment of a disease condition associated with a DAM is described. The
ScFv Ab has therapeutic, diagnostic and prognostic applications.
ICM C07K016-00
15-3 (Immunochemistry)
Section cross-reference(s): 3, 63
chimeric antibody tumor assocd antigen antitumor; scFv disease
assocd mol diagnosis prognosis
Immunoglobulins
RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP
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Immunoglobulins
RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
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   (G; chimeric antibody scFv recognizing disease-associated mol.
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Immunostimulants
   (adjuvants; chimeric antibody scFv recognizing
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   (cancer; chimeric antibody scFv recognizing disease-associated
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Antitumor agents
Bacteriophage
Carcinoma
DNA sequences
Diagnosis
Disease, animal
Drugs
Genetic vectors
Human immunodeficiency virus
Imaging
Molecular cloning
Plasmids
Prognosis
Protein sequences
Protein sequences
   (chimeric antibody scFv recognizing disease-associated mol. for
   diagnosis and treatment of disease)
Antibodies
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Fusion proteins (chimeric proteins)

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Proteins, general, biological studies
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (chimeric antibody scFv recognizing disease-associated mol. for
        diagnosis and treatment of disease)
IT
     Promoter (genetic element)
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES
     (Uses)
        (chimeric antibody scFv recognizing disease-associated mol. for
        diagnosis and treatment of disease)
TT
     CD28 (antigen)
     RL: BPR (Biological process); BSU (Biological study, unclassified); THU
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        (chimeric antibody scFv recognizing disease-associated mol. for
        diagnosis and treatment of disease)
IT
     CTLA-4 (antigen)
     RL: BPR (Biological process); BSU (Biological study, unclassified); THU
     (Therapeutic use); BIOL (Biological study); PROC (Process); USES (Uses)
        (chimeric antibody scFv recognizing disease-associated mol. for
        diagnosis and treatment of disease)
IT
     Antigens
     Toxins
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     (Biological study); USES (Uses)
        (chimeric antibody scFv recognizing disease-associated mol. for
        diagnosis and treatment of disease)
IT
     CD80 (antigen)
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        (chimeric; chimeric antibody scFv recognizing
        disease-associated mol. for diagnosis and treatment of disease)
ΙT
     Neoplasm
        (diagnosis; chimeric antibody scFv recognizing
        disease-associated mol. for diagnosis and treatment of disease)
IT
     Immunoglobulins
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
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     (Preparation); USES (Uses)
        (fragments, scFv; chimeric antibody scFv recognizing
        disease-associated mol. for diagnosis and treatment of disease)
TT
     Interleukin 5
     RL: BPN (Biosynthetic preparation); PRP (Properties); THU (Therapeutic
     use); BIOL (Biological study); PREP (Preparation); USES (Uses)
        (fusion protein; chimeric antibody scFv recognizing
        disease-associated mol. for diagnosis and treatment of disease)
IT
     Envelope proteins
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (qp120env; chimeric antibody scFv recognizing disease-associated
        mol. for diagnosis and treatment of disease)
IT
     Immunoglobulins
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (heavy chains; chimeric antibody scFv recognizing
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disease-associated mol. for diagnosis and treatment of disease)
     Enzymes, biological studies
IT
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES
        (pro-drug activating; chimeric antibody scFv recognizing
        disease-associated mol. for diagnosis and treatment of disease)
IT
     Drug delivery systems
        (prodrugs; chimeric antibody scFv recognizing disease-associated
        mol. for diagnosis and treatment of disease)
IT
     Antigens
     RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
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     Antigens
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L16 ANSWER 36 OF 125 CAPLUS COPYRIGHT 2006 ACS on STN
                        2001:168142 CAPLUS
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                         134:232695
DOCUMENT NUMBER:
TITLE:
                         Nucleic acids encoding human secreted and
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                         Eaton, Dan L.; Filvaroff, Ellen; Gerritsen, Mary E.;
INVENTOR(S):
                         Goddard, Audrey; Godowski, Paul J.; Grimaldi,
                         Christopher J.; Gurney, Austin L.; Watanabe, Colin K.;
                         Wood, William I.
PATENT ASSIGNEE(S):
                         Genentech, Inc., USA
                         PCT Int. Appl., 278 pp.
SOURCE:
                         CODEN: PIXXD2
DOCUMENT TYPE:
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                         English
LANGUAGE:
FAMILY ACC. NUM. COUNT: 143
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US 2000-201516P
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WO 2000-US13358
                    Α
                       20000515
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ED Entered STN: 09 Mar 2001

The present invention is directed to novel human polypeptides and to AB nucleic acid mols. encoding those polypeptides. Eighty-four cDNAs clones were identified by homol. screening of extracellular domains (including the secretion signal sequence, if any), amylase screening, and signal algorithm anal. The proteins possess various useful biol. activities: including (1), pericyte c-fos induction, (2) affecting the release of glucose or free fatty acid uptake in skeletal muscle, (3) stimulating the release of proteoglycans from cartilage, and (4) stimulating tumor necrosis factor- α release in human blood. Many of the proteins show differential expression between normal and tumor tissues. Also provided herein are vectors and host cells comprising those nucleic acid sequences, chimeric polypeptide mols. comprising the polypeptides of the present invention fused to heterologous polypeptide sequences, antibodies which bind to the polypeptides of the present invention and to methods for producing the polypeptides of the present invention.

IC ICM C12N015-12

ICS C07K014-47; C07K014-705; G01N033-53; C12N015-62; C07K016-18

CC 3-3 (Biochemical Genetics)

Section cross-reference(s): 6, 13, 63

IT Immunoglobulins

RL: BPN (Biosynthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(fusion products; nucleic acids encoding human secreted and transmembrane polypeptides)

IT Epitopes

(fusion proteins containing; nucleic acids encoding human secreted and transmembrane polypeptides)

IT Fusion proteins (chimeric proteins)

RL: BPN (Biosynthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(with epitope tags or Fc region of Ig; nucleic acids encoding human secreted and transmembrane polypeptides)

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221877-81-2P
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225371-37-9P
               229324-31-6P, Interleukin 1δ (mouse)
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243122-70-5P
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IT

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     (Biological study, unclassified); PRP (Properties); THU (Therapeutic use);
    BIOL (Biological study); OCCU (Occurrence); PREP (Preparation); USES
     (Uses)
        (amino acid sequence; nucleic acids encoding human secreted and
       transmembrane polypeptides)
L16 ANSWER 37 OF 125 CAPLUS COPYRIGHT 2006 ACS on STN
                        2000:911441 CAPLUS
ACCESSION NUMBER:
                        134:68048
DOCUMENT NUMBER:
                        Analogs of a cytochrome P450 of Pseudomonas putida
TITLE:
                        with improved catalytic action aromatic
                        halohydrocarbons for use in bioremediation of soil
                        Wong, Luet Lok; Jones, Jonathan Peter
INVENTOR(S):
                        Isis Innovation Limited, UK
PATENT ASSIGNEE(S):
                        PCT Int. Appl., 42 pp.
SOURCE:
                        CODEN: PIXXD2
                        Patent
DOCUMENT TYPE:
                        English
LANGUAGE:
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
                                         APPLICATION NO.
                       KIND DATE
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PRIORITY APPLN. INFO.:
                                                             W 20000619
                                           WO 2000-GB2379
     Entered STN: 29 Dec 2000
     Analaogs of cytochrome P 450cam of Pseudomonas putida that have improved
     catalytic activity against heavily halogenated aromatic hydrocarbons and that
     may be of use in the reclamation of soils contaminated with
     polychlorinated biphenyls. In particular, alterations in the substrate
     pocket that increase the volume available for bulky polyhalogenated aroms.
     are described. Preparation of a series of analogs of the gene camC cytochrome
     P 450 with increased activity towards polychlorinated biphenyls is
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demonstrated. A fusion protein of putidaredoxin and putidaredoxin

reductase that can be used as a cofactor is also described.

ED

AB

IC

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CC
    7-5 (Enzymes)
    Section cross-reference(s): 3, 10, 19, 60
IT
    Ferredoxins
    RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL
     (Biological study); PROC (Process)
        (putidaredoxins, cofactors for cytochrome P 450, fusion
       products; analogs of cytochrome P 450 of Pseudomonas putida with
        improved catalytic action aromatic halohydrocarbons for use in
       bioremediation of soil)
    9059-45-4D, Putidaredoxin reductase, fusion products with
IT
    putidaredoxin
    RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL
     (Biological study); PROC (Process)
        (cofactor for cytochrome P 450; analogs of cytochrome P 450 of
        Pseudomonas putida with improved catalytic action aromatic
       halohydrocarbons for use in bioremediation of soil)
    315722-77-1
IT
    RL: PRP (Properties)
        (unclaimed protein sequence; analogs of a cytochrome P 450 of
        Pseudomonas putida with improved catalytic action aromatic
       halohydrocarbons for use in bioremediation of soil)
REFERENCE COUNT:
                              THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS
                        7
                              RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
L16 ANSWER 38 OF 125 CAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                        2000:900835 CAPLUS
DOCUMENT NUMBER:
                        134:52293
TITLE:
                        Human and murine secreted or transmembrane proteins
                        and their encoding nucleic acids having diagnostic,
                        preventive, therapeutic, and other uses
INVENTOR(S):
                        Mccarthy, Sean A.; Fraser, Christopher C.; Sharp, John
                        D.; Barnes, Thomas M.
PATENT ASSIGNEE(S):
                        Millennium Pharmaceuticals, Inc., USA
SOURCE:
                        PCT Int. Appl., 359 pp.
                        CODEN: PIXXD2
DOCUMENT TYPE:
                        Patent
LANGUAGE:
                        English
FAMILY ACC. NUM. COUNT:
                        10
PATENT INFORMATION:
                                        APPLICATION NO.
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                       KIND DATE
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            LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD,
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            IE, SI, LT, LV, FI, RO, MK, CY, AL
PRIORITY APPLN. INFO.:
                                           US 1999-333159
                                                              A 19990614
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ED Entered STN: 22 Dec 2000

AB The invention provides isolated nucleic acids encoding a variety of

WO 2000-US14858

W 20000524

proteins having diagnostic, preventive, therapeutic, and other uses. Thus, the sequences for 6 human and 2 murine cDNA mols. are provided for proteins designated TANGO 202, TANGO 265, TANGO 273, TANGO 286, TANGO 294, and INTERCEPT 296. Tissue distribution, biol. functions, and chromosomal gene mapping are also provided. These nucleic acids and proteins are useful for diagnosis, prevention, and therapy of a number of human and other animal disorders. The invention also provides antisense nucleic acid mols., expression vectors containing the nucleic acid mols. of the invention, host cells into which the expression vectors have been introduced, and non-human transgenic animals in which a nucleic acid mol. of the invention has been introduced or disrupted. The invention still further provides isolated polypeptides, fusion polypeptides, antiqenic peptides and antibodies. Diagnostic, screening, and therapeutic methods utilizing compns. of the invention are also provided. The nucleic acids and polypeptides of the present invention are useful as modulating agents in regulating a variety of cellular processes.

TC ICM C12Q

CC 3-3 (Biochemical Genetics)

Section cross-reference(s): 6, 13, 63

TT Antibodies

Fusion proteins (chimeric proteins)

Primers (nucleic acid)

Probes (nucleic acid)

RL: BPN (Biosynthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(human and murine secreted or transmembrane proteins and their encoding nucleic acids having diagnostic, preventive, therapeutic, and other uses)

IT 143891-11-6 149223-58-5 154400-02-9, Protein (Caenorhabditis elegans clone C06E1 gene C06E1.3 reduced) 168458-95-5, Semaphorin SemB 313413-97-7 313413-98-8 313413-99-9 313414-00-5 313414-01-6 313414-02-7 313414-03-8 313414-04-9 313414-05-0 313414-06-1 313414-07-2 313484-72-9 313484-73-0 313484-74-1 313484-75-2 313484-76-3 313484-77-4 RL: PRP (Properties)

(unclaimed protein sequence; human and murine secreted or transmembrane proteins and their encoding nucleic acids having diagnostic, preventive, therapeutic, and other uses)

L16 ANSWER 39 OF 125 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2000:900777 CAPLUS

DOCUMENT NUMBER: 134:37967

TITLE: Cloning and cDNA and deduced amino acid sequences of

42 human secreted proteins

INVENTOR(S): Rosen, Craig A.; Ruben, Steven M.; Komatsoulis, George

Α.

PATENT ASSIGNEE(S): Human Genome Sciences, Inc., USA

SOURCE: PCT Int. Appl., 453 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

English LANGUAGE:

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.						KIND		DATE			APPLICATION NO.					DATE			
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             AZ, BY, KG, KZ, MD, RU, TJ, TM
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     EP 1187908
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            AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
         R:
             IE, SI, LT, LV, FI, RO
                                20030121
     JP 2003502031
                          T2
                                            JP 2001-503618
                                                                    20000601
                                            US 1999-138573P
                                                                 Р
                                                                    19990611
PRIORITY APPLN. INFO.:
                                            US 2000-174851P
                                                                 P
                                                                    20000107
                                            WO 2000-US14929
                                                                 W
                                                                    20000601
ED
     Entered STN: 22 Dec 2000
     The present invention relates to 42 novel human secreted proteins and
AB
     isolated nucleic acids containing the coding regions of the genes encoding
     such proteins. Tissue distribution, sequence homologies, and preferred
     epitope sites are provided for the secreted proteins, as well as
     chromosomal mapping of some of the genes. Also provided are vectors, host
     cells, antibodies, and recombinant methods for producing human secreted
     proteins in bacterial, insect, and mammalian cells. The invention further
     relates to diagnostic and therapeutic methods useful for diagnosing and
     treating disorders related to these novel human secreted proteins.
     High-throughput screening assays are also provided for various putative
     activities (no data) of the secreted proteins.
     ICM C12N001-21
TC:
     ICS
         C12N005-10; C12N015-12; C12N015-63; C07K014-435
CC
     3-3 (Biochemical Genetics)
     Section cross-reference(s): 6, 13, 63
     Immunoglobulins
IT
     RL: BPN (Biosynthetic preparation); THU (Therapeutic use); BIOL
     (Biological study); PREP (Preparation); USES (Uses)
        (fusion products; cloning and cDNA and deduced amino acid
        sequences of 42 human secreted proteins)
                    312779-61-6P
                                   312779-62-7P 312779-63-8P
IT
     312779-60-5P
                    312779-65-0P
                                   312779-66-1P
                                                  312779-67-2P
                                                                  312909-32-3P
     312779-64-9P
                    312909-34-5P
                                   312909-35-6P
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     312909-33-4P
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     312909-38-9P
                    312909-39-0P
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     312909-48-1P
                                                  312909-51-6P
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     312909-53-8P
                    312909-56-1P
                                   312909-58-3P
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                    312909-62-9P
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     312909-66-3P
                    312909-73-2P
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     312909-72-1P
     RL: BOC (Biological occurrence); BPN (Biosynthetic preparation); BSU
     (Biological study, unclassified); PRP (Properties); THU (Therapeutic use);
     BIOL (Biological study); OCCU (Occurrence); PREP (Preparation); USES
     (Uses)
        (amino acid sequence; cloning and cDNA and deduced amino acid sequences
        of 42 human secreted proteins)
     312781-77-4
                   312781-78-5
                                 312781-79-6
                                                312781-80-9
                                                              312781-81-0
IT
                                 312781-84-3
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     312781-82-1
                   312781-83-2
     312781-87-6
                   312781-88-7 312781-89-8
                                            312781-90-1
     312781-91-2
                   312909-31-2
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                                                              312909-82-3
                                 312955-26-3
                                               312955-27-4
     312909-83-4
                   312909-92-5
     RL: PRP (Properties)
        (unclaimed protein sequence; cloning and cDNA and deduced amino acid
        sequences of 42 human secreted proteins)
                               THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS
REFERENCE COUNT:
                         3
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RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

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L16 ANSWER 40 OF 125 CAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                        2000:881193 CAPLUS
DOCUMENT NUMBER:
                        134:51375
                        Modulation of protein levels using the SCF complex
TITLE:
                        Zhang, Hui; Tsvetkov, Lyuben M.; Kondo, Takeshi
INVENTOR(S):
                        Yale University, USA
PATENT ASSIGNEE(S):
                        PCT Int. Appl., 162 pp.
SOURCE:
                        CODEN: PIXXD2
DOCUMENT TYPE:
                        Patent
                        English
LANGUAGE:
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
                                          APPLICATION NO.
                       KIND DATE
    PATENT NO.
     _____
                               _____
                       - - - <del>-</del>
                        A1 20001214 WO 2000-US15449
                                                                20000605 <--
    WO 2000075184
        W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR,
            CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU,
             ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU,
             LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE,
            SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA,
             ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
             DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ,
             CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
PRIORITY APPLN. INFO.:
                                          US 1999-137494P
     Entered STN: 15 Dec 2000
     This invention encompasses various methods of modulating protein levels
AB
     using the SKP1, CDC53/Cullin, F-box(SCF) protein complex. More
     specifically, the present invention provides various methods of target
     protein degradation using targeted ubiquitination techniques. The present
     invention also provides various compns. and assays associated with the
     disclosed modulation of protein levels using the SCF complex as well as
     various methods of detecting, monitoring and treating cancerous cells.
     ICM C07K014-47
IC
     ICS A61K038-17; G01N033-68; A61P043-00
CC
     1-6 (Pharmacology)
     Section cross-reference(s): 3, 9
     Gene, animal
IT
     RL: BAC (Biological activity or effector, except adverse); BPR (Biological
     process); BSU (Biological study, unclassified); BIOL (Biological study);
     PROC (Process)
        (SKP2-target protein interaction domain-containing protein fusion
        protein-encoding; modulation of protein levels using SCF
        (SKP1-CDC53/Cullin-F-box) complex in relation to targeted
        ubiquitination techniques and cancer monitoring and treatment)
     Fusion proteins (chimeric proteins)
IT
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); BIOL (Biological study)
        (of SKP2 and target protein interaction domain-containing protein.;
        modulation of protein levels using SCF (SKP1-CDC53/Cullin-F-box)
        complex in relation to targeted ubiquitination techniques and cancer
        monitoring and treatment)
     Proteins, specific or class
IT
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); BIOL (Biological study)
        (target protein interaction domain-containing, fusion protein
        with SKP2; modulation of protein levels using SCF (SKP1-CDC53/Cullin-F-
        box) complex in relation to targeted ubiquitination techniques and
```

cancer monitoring and treatment)

IT 85256-55-9, Phosphoprotein (human gene c-myc protein moiety reduced) 143861-61-4, Cyclin E (human) 147277-84-7, Protein (human clone C1 gene max 104-amino acid isoform) 147572-23-4, Protein (mouse clone 1 147707-23-1, Protein (human gene MDM2) 422-amino acid reduced) 148846-23-5, Protein (human clone Mad-1 gene mad reduced) 151440-09-4, Protein (mouse RL-7 cell gene bax isoform α reduced) 157908-85-5 183627-31-8, Protein (human gene BCL-2) 202220-27-7 170618-97-0 208541-48-4, Protein p53 (human) 207624-75-7 222963-35-1 251650-48-3 251650-57-4 251654-20-3 251654-24-7 260421-71-4 313408-60-5, Protein (human gene ZF1) 313408-61-6, Protein (human gene ZF3) 313408-62-7, Protein (mouse gene ZF5) 313408-63-8, Protein (human gene 313408-64-9, Protein (human gene ZF9) 313408-65-0, Protein (mouse 313408-66-1, Protein (human gene ZF19) 313408-67-2, Protein gene ZF13) (mouse gene ZF23) 313408-68-3, Protein (human gene ZF25) RL: BOC (Biological occurrence); BPR (Biological process); BSU (Biological study, unclassified); PRP (Properties); BIOL (Biological study); OCCU (Occurrence); PROC (Process)

(amino acid sequence; modulation of protein levels using SCF (SKP1-CDC53/Cullin-F-box) complex in relation to targeted ubiquitination techniques and cancer monitoring and treatment)

REFERENCE COUNT: 13 THERE ARE 13 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L16 ANSWER 41 OF 125 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2000:875755 CAPLUS

DOCUMENT NUMBER: 134:41092

TITLE: Compounds and methods for treatment and diagnosis of

mycobacterial infections

INVENTOR(S): Visser, Elizabeth

PATENT ASSIGNEE(S): Genesis Research and Development Corporation Limited,

N. Z.

SOURCE: U.S., 147 pp., Cont.-in-part of U.S. 5,985,287.

CODEN: USXXAM

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 8

PATENT INFORMATION:

PA.	CENT :	NO.			KIN	D	DATE			APPL	ICAT	ION I	NO.			ATE		
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US	6160	093			Α		2000	1212		US 1	998-	9585	5		1:	9980	611 <	<
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US	5985	287			A		1999	1116		US 1	997-	9973	62		1	9971:	223 4	<
	6406						2002						-					
				20020618 US 1998-205426 19990701 CA 1998-2315539														
					AA		1999	0/01										
WO	9932	932634 A2				19990701			WO 1998-NZ189					19981223 <				
WO	9932	634			A 3		1999	1202										
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		ТJ,	TM															
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ΑU	9918	936			A1		1999	0712		AU 1	999-	1893	6		1:	9981	223 <	<
ΑU	7463	11			B2		2002	0418										

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BR 1998-14432
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                               20001010
                                           EP 1998-963665
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                               20001018
    EP 1044273
                         A2
            AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
        R:
             IE, FI
                                           TR 2000-200001948
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                               20010221
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                               20021026
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                                           US 1996-705347
                                                               A2 19960829
PRIORITY APPLN. INFO.:
                                                               A2 19970612
                                           US 1997-873970
                                                               A2 19971223
                                           US 1997-997362
                                           US 1997-996624
                                                               A 19971223
                                                               A 19971223
                                           US 1997-997080
                                            IN 1998-CA242
                                                               A 19980216
                                           US 1998-95855
                                                               A2 19980611
                                           US 1998-156181
                                                               A 19980917
                                                               A 19981204
                                           US 1998-205426
                                                               W 19981223
                                           WO 1998-NZ189
     Entered STN: 14 Dec 2000
ED
     The present invention provides polypeptides comprising an immunogenic
AB
    portion of a M. vaccae protein and DNA mols. encoding such polypeptides,
     together with methods for their use in the diagnosis and treatment of
     mycobacterial infection. Methods for enhancing the immune response to an
     antigen including administration of M. vaccae culture filtrate,
     delipidated M. vaccae cells or delipidated and deglycolipidated M. vaccae
     cells are also provided.
IC
     ICM C07K014-35
     ICS A61K039-04
INCL 530350000
     15-2 (Immunochemistry)
CC
     Section cross-reference(s): 3, 9
     Fusion proteins (chimeric proteins)
TT
     RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (Mycobacterium vaccae antigens as vaccine, diagnostic,
        immunotherapeutic and adjuvant for infection, immunol. disease, and
        cancer)
                   204785-66-0, Antigen 85B (Mycobacterium vaccae fragment)
     204785-65-9
IT
                   204785-82-0 204785-84-2, Antigen GV-24 (Mycobacterium
     204785-67-1
              204785-86-4, Antigen GV-25 (Mycobacterium vaccae)
     204785-97-7, Antigen GV-27 (Mycobacterium vaccae N-terminal
     fragment) 204786-00-5, Antigen GV-27A (Mycobacterium vaccae
     N-terminal fragment)
                           204786-01-6, Antigen GV-27B (Mycobacterium vaccae
                228106-68-1, Antigen GV-22B (Mycobacterium vaccae)
     228107-02-6, Antigen GV-1/83 (Mycobacterium vaccae)
                                                          228107-35-5, Antigen
     GVs-9 (Mycobacterium vaccae) 228107-42-4, Antigen GV-27
                             228107-43-5 228107-53-7, Antigen GV-24B
     (Mycobacterium vaccae)
     (Mycobacterium vaccae)
                              228107-62-8, Antigen GV-38A (Mycobacterium
               228107-65-1, Antigen GV-38B (Mycobacterium vaccae)
     228107-79-7, Antigen GV-33 (Mycobacterium vaccae)
                                                        228107-83-3, Antigen
     GVc-13 (Mycobacterium vaccae)
                                   228107-96-8, Antigen GVs-9 (Mycobacterium
               228107-99-1, Antigen GV-29 (Mycobacterium vaccae)
     Antigen GV-45 (Mycobacterium vaccae)
                                          228108-08-5, Antigen GV-41B
     (Mycobacterium vaccae) 228108-12-1, Antigen GV-44 (Mycobacterium vaccae)
                 250277-71-5
     228108-14-3
     RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
     (Biological study)
        (amino acid sequence; Mycobacterium vaccae antigens as vaccine,
        diagnostic, immunotherapeutic and adjuvant for infection, immunol.
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disease, and cancer)

119939-22-9, Antigen α (Mycobacterium BCG clone p $\alpha L-1$ IT136108-01-5, Antigen (Mycobacterium tuberculosis precursor reduced) 32.0-kilodalton precursor reduced) 145110-35-6, Antigen 85C (Mycobacterium tuberculosis clone 11-2 precursor) 146045-21-8, Antigen 85B (Mycobacterium leprae precursor reduced) 152206-84-3 157908-64-0 204785-69-3, Antigen GVc-7 (Mycobacterium vaccae) 204785-74-0 204785-78-4 204785-75-1 204785-76-2 204785-77-3 204785-80-8 204785-88-6 204785-93-3 204785-95-5 204786-03-8 204786-06-1 204786-07-2 204786-10-7 204786-11-8 204786-15-2 204786-16-3 228105-87-1, Antigen GVc-13 (Mycobacterium vaccae) 228107-33-3, Antigen GV-1/70 (Mycobacterium vaccae) 228107-40-2, Antigen GV-5P (Mycobacterium 228107-44-6, Antigen GV-27B (Mycobacterium vaccae) 228107-47-9 228107-48-0 228107-68-4, Antigen GV-41 (Mycobacterium vaccae) 228107-70-8, Antigen GV-42 (Mycobacterium vaccae) 228107-73-1, Antigen GV-44 (Mycobacterium vaccae) 228107-76-4 228111-24-8 228401-90-9 250261-47-3 250277-23-7 **250277-25-9** 250277-35-1 250277-56-6 313050-89-4 313050-94-1 313050-97-4 313050-99-6 313051-00-2 313051-40-0 313051-41-1 313051-42-2 313051-43-3 313051-44-4 313051-45-5 313051-46-6 313051-47-7 313051-59-1 313051-69-3 313051-60-4 313051-61-5 313051-62-6 313051-63-7 313051-70-6 313051-71-7 313051-72-8 313051-73-9 313051-74-0 313051-88-6 313051-89-7 313051-90-0 313051-91-1 313051-93-3 RL: PRP (Properties)

(unclaimed protein sequence; compds. and methods for treatment and diagnosis of mycobacterial infections)

REFERENCE COUNT: 19 THERE ARE 19 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L16 ANSWER 42 OF 125 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2000:874126 CAPLUS

DOCUMENT NUMBER: 134:41111

TITLE: Viral vectors to inhibit leukocyte infiltration or

cartilage degradation of joints

INVENTOR(S): Glorioso, Joseph C.; Evans, Christopher H.; Robbins,

Paul D.; Ghivizzani, Steven C.

PATENT ASSIGNEE(S): University of Pittsburgh of the Commonwealth System of

Higher Education, USA

SOURCE: U.S., 72 pp., Cont.-in-part of U.S. Ser. No. 685,212.

CODEN: USXXAM

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 8

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 6159464	Α	20001212	US 1997-924376	19970905 <
US 6228356	B1	20010508	US 1996-685212	19960723
US 2003220283	A1	20031127	US 2003-366123	20030212
PRIORITY APPLN. INFO.:			US 1990-630981 B:	2 19901220
			US 1993-27750 B	1 19930308
			US 1996-685212 A	2 19960723
			US 1992-963928 B	2 19921020
			US 1997-924376 A	1 19970905
			US 2000-626597 B:	1 20000727

ED Entered STN: 14 Dec 2000

AB Methods for treating a connective tissue disorder by introducing at least one gene encoding a product into at least one target cell of a mammalian host for use in treating the mammalian host are disclosed. These methods

include employing recombinant techniques to produce a vector mol. containing the DNA sequence encoding for the product and infecting the target cell of the mammalian host using the vector. The injection can be done in vivo, by directly injecting the vector into the host, or can be done in vitro by transfecting a population of cultured target cells with the vector and transplanting them each into the host. Nonviral means can also be used to introduce the DNA sequence to the host. Administration of more than one gene of interest results in an enhanced therapeutic benefit. Also disclosed is a method for treating a connective tissue disorder by introducing at least one gene encoding a product into at least one target cell of a joint of a host for use in treating multiple joints of the host. Injection of a vector mol. containing the DNA sequence encoding for a product of interest, or non-viral introduction of such a DNA sequence, to one join of a mammalian host results in a therapeutic benefit in that joint as well as other joints in the host. ICM A01N063-00

IC ICS A61K048-00

INCL 424093200

15-5 (Immunochemistry)

Section cross-reference(s): 3, 63

IT Immunoglobulins

> RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(fusion products; viral vectors encoding human interleukin 1 receptor antagonist or related protein for gene therapy of connective tissue disorders)

ΙT Fusion proteins (chimeric proteins)

Gene, microbial Interleukin 10 Interleukin 1a Interleukin 1B

Promoter (genetic element)

Tumor necrosis factors

RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(viral vectors encoding human interleukin 1 receptor antagonist or related protein for gene therapy of connective tissue disorders)

118901-67-0 124541-29-3 IT

> RL: PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(amino acid sequence; viral vectors encoding human interleukin 1 receptor antagonist or related protein for gene therapy of connective tissue disorders)

REFERENCE COUNT: 27 THERE ARE 27 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L16 ANSWER 43 OF 125 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2000:861924 CAPLUS

DOCUMENT NUMBER: 134:40682

TITLE: Breast, gastric and prostate cancer-associated antigens and their diagnostic and therapeutic uses

INVENTOR(S): Obata, Yuichi

Ludwig Institute for Cancer Research, USA PATENT ASSIGNEE(S):

SOURCE: PCT Int. Appl., 799 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

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PATENT NO.
                        KIND
                               DATE
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                                                                  _____
                                           WO 2000-US14749
     WO 2000073801
                        A2
                               20001207
                                                                  20000526 <--
     WO 2000073801
                         А3
                               20020912
        W: AU, CA, CN, JP, KR, US
        RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL,
            PT. SE
    EP 1259812
                         A2
                               20021127
                                           EP 2000-932804
                                                                  20000526
            AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
        R:
             IE, FI, CY
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                               20030610
                                           JP 2001-500870
     JP 2003518364
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PRIORITY APPLN. INFO.:
                                           US 1999-136526P
                                                               P 19990528
                                           US 1999-153454P
                                                               P 19990910
                                           WO 2000-US14749
                                                               W
                                                                  20000526
     Entered STN: 08 Dec 2000
ED
AB
     Cancer-associated antigens have been identified by autologous antibody
     screening of libraries of nucleic acids expressed in breast, gastric, and
    prostate cancer cells using antisera from cancer patients. The invention
    relates to 593 nucleic acids and 740 encoded polypeptides which are
     cancer-associated antigens expressed in patients afflicted with cancer.
     invention provides, inter alia, isolated nucleic acid mols., expression
     vectors containing those mols., and host cells transfected with those mols.
     The invention also provides isolated proteins and peptides, antibodies to
     those proteins and peptides and cytotoxic T lymphocytes which recognize
     the proteins and peptides. Fragments of the foregoing including
     functional fragments and variants also are provided. Kits containing the
     foregoing mols. addnl. are provided. The mols. provided by the invention
     can be used in the diagnosis, monitoring, research, or treatment of
     conditions characterized by the expression of one or more cancer associated
     antigens.
IC
     ICM G01N033-574
         C12Q001-68; A61K039-00; A61K039-395; A61K035-14; A61K048-00;
     ICS
          C12N015-63; C12N005-10; C07K014-47; C07K007-04; C07K016-18
     14-1 (Mammalian Pathological Biochemistry)
CC
     Section cross-reference(s): 3, 9, 15, 63
IT
    Antibodies
     RL: BPR (Biological process); BSU (Biological study, unclassified); THU
     (Therapeutic use); BIOL (Biological study); PROC (Process); USES (Uses)
        (chimeric; breast, gastric and prostate cancer-associated
        antigens and their diagnostic and therapeutic uses)
                   312642-68-5P
IT
     312642-67-4P
                                  312642-69-6P
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RL: ANT (Analyte); BOC (Biological occurrence); BPN (Biosynthetic preparation); BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); OCCU (Occurrence); PREP (Preparation); USES (Uses)

(amino acid sequence; breast, gastric and prostate cancer-associated antigens and their diagnostic and therapeutic uses)

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RL: ANT (Analyte); BOC (Biological occurrence); BPN (Biosynthetic preparation); BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); OCCU (Occurrence); PREP (Preparation); USES (Uses)

(amino acid sequence; breast, gastric and prostate cancer-associated antigens and their diagnostic and therapeutic uses)

L16 ANSWER 44 OF 125 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2000:861814 CAPLUS

DOCUMENT NUMBER: 134:26782

TITLE: Identification and characterization of Snf2 related

CBP activator protein (SRCAP)

INVENTOR(S): Chrivia, John; Yaciuk, Peter PATENT ASSIGNEE(S): Saint Louis University, USA SOURCE: PCT Int. Appl., 103 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.				KIND DATE		APPLICATION NO.					DATE						
						-									_		
WO	2000	0734	67		A1		2000	1207	1	WO 2	000-1	JS14	719		2	0000!	525 <
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US	6365	372			В1		2002	0402	1	JS 2	000-	5791	31		2	0000	525
PRIORIT	Y APP	LN.	INFO	. :					1	JS 19	999-	1366:	20P]	P 19	9990	527
									1	JS 20	000-	5791	31	i	A 20	0000	525
			•			^ ^											

ED Entered STN: 08 Dec 2000

AB A protein, SRCAP, a novel SNF2/SWI2 protein family member interacting with CREB-binding proteins, is provided. The protein is capable of co-activating CREB binding protein (CBP) mediated transcription, as well

as activating transcription without CBP. SRCAP is a Snf2 family member. As such, it has ATPase activity. Consistent with this activity, SRCAP contains the conserved ATPase domain found within members of the Snf2 family. Transfection expts. demonstrate that SRCAP is able to activate transcription when expressed as a Gal-SRCAP chimera and that SRCAP also enhances the ability of CBP to activate transcription. The adenoviral protein E1A is found to disrupt interaction between SRCAP and CBP possibly representing a mechanism for E1A-mediated transcriptional repression. SRCAP also interacts with NS5A of hepatitis C virus (HCV), and this interaction may have effect on growth regulation of cells infected with HCV through the down-regulation of p21 promoter activity and contribute HCV pathogenesis. Fragments of SRCAP are also provided, as are its cDNA and cDNA fragments. Antibodies that bind to SRCAP are also provided.

IC ICM C12N015-52

ICS C12N015-62; C12N015-11; C12N005-10; C12N009-14; C07K016-40; G01N033-50; G01N033-566; C12Q001-68; A61K038-46

CC 6-3 (General Biochemistry)

Section cross-reference(s): 3, 13

IT Transcription factors

RL: BPR (Biological process); BSU (Biological study, unclassified); BUU (Biological use, unclassified); BIOL (Biological study); PROC (Process); USES (Uses)

(GAL4, fusion product with SRCAP; cloning and sequence of transcription factor SRCAP which interacts with CREB-binding protein)
Fusion proteins (chimeric proteins)

RL: BPR (Biological process); BSU (Biological study, unclassified); BUU (Biological use, unclassified); BIOL (Biological study); PROC (Process); USES (Uses)

(of GAL4 and SRCAP; cloning and sequence of transcription factor SRCAP which interacts with CREB-binding protein)

IT 311824-07-4 311824-09-6

RL: BPR (Biological process); BSU (Biological study, unclassified); PRP (Properties); BIOL (Biological study); PROC (Process)

(amino acid sequence; cloning and sequence of transcription factor SRCAP which interacts with CREB-binding protein)

REFERENCE COUNT:

IT

THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L16 ANSWER 45 OF 125 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2000:861811 CAPLUS

DOCUMENT NUMBER: 134:26781

TITLE: Chaperonin GroEL muteins with improved stability

INVENTOR(S): Buckle, Ashley Maurice; Fersht, Alan

PATENT ASSIGNEE(S): Medical Research Council, UK

SOURCE: PCT Int. Appl., 58 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND DATE	APPLICATION NO.	DATE		
WO 2000073463	A1 20001207	WO 2000-GB2019	20000526 <		
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LV, MA, MD	, MG, MK, MN, MW,	MX, MZ, NO, NZ, PL, PT,	RO, RU, SD,		
SE, SG, SI	, SK, SL, TJ, TM,	TR, TT, TZ, UA, UG, US,	UZ, VN, YU,		
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PRIORITY APPLN. INFO.:
                                            GB 1999-12445
                                                                  19990527
                                            WO 2000-GB2019
                                                                  20000526
ED
     Entered STN: 08 Dec 2000
AB
     A GroEL chaperone polypeptide, or homolog thereof, or fragment thereof
     having protein refolding activity, comprising one or more amino acid
     modifications at any one of amino acid residues (207, 212, 217, 223, 233,
     267, 271, 294, 305, 308 and 326) of the Escherichia coli GroEL amino acid
     sequence or their equivalent positions in other homologous chaperone
     polypeptides is provided. Sequence comparison of Escherichia coli
     GroEL(193-345) with the apical domains of 129 homologous Cpn60 proteins
     available in the public sequence databanks reveals that 31 of the E. coli
     GroEL(193-345) residues occur with a frequency of less than 35%. These
     amino acid residues were were substituted in the GroEL sequence by the
     dominant residues at the resp. positions with the aim of obtaining more
     stable variants. At least 13 substitutions are identified that increase
     protein stability. By combining the stabilizing mutations, two multiple
     mutants were created that have significantly increased stability while
     retaining full chaperone activity: A212E/A223T/M223L/I305L/E308K/N326T and
     A212E/A223V/M233L/I305L/E308K/N326T. The stabilized GroEL muteins may be
     used for reconditioning of other mols. (e.g., in vitro refolding of human
     cyclophilin A) after they have been inactivated or denatured.
IC
     ICM C12N015-31
     ICS C07K014-245; C12N015-62; A61K038-16; C07K016-12
     6-3 (General Biochemistry)
CC
     Section cross-reference(s): 3, 9, 63
IT
     Fusion proteins (chimeric proteins)
     RL: BPN (Biosynthetic preparation); BUU (Biological use, unclassified);
     THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES
     (Uses)
        (chaperonin GroEL muteins with improved stability)
IT
     169444-45-5DP, GroEL chaperonin (Escherichia coli strain
     K12-MG1655 gene mopA/groEL), muteins 205541-08-8DP, muteins
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     RL: BPN (Biosynthetic preparation); BUU (Biological use, unclassified);
     PEP (Physical, engineering or chemical process); PRP (Properties); BIOL
     (Biological study); PREP (Preparation); PROC (Process); USES (Uses)
        (amino acid sequence; chaperonin GroEL muteins with improved stability)
                               THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS
REFERENCE COUNT:
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                               RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
L16 ANSWER 46 OF 125 CAPLUS COPYRIGHT 2006 ACS on STN
                        2000:772768 CAPLUS
ACCESSION NUMBER:
DOCUMENT NUMBER:
                        133:334034
TITLE:
                        VlsE-derived peptides, nucleic acids, and vaccines for
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diagnosis and prevention of Lyme disease

Philipp, Mario T.; Liang, Fang Ting INVENTOR (S): The Administrators of the Tulane Educational Fund, USA PATENT ASSIGNEE(S): PCT Int. Appl., 76 pp. SOURCE: CODEN: PIXXD2 DOCUMENT TYPE: Patent English LANGUAGE: FAMILY ACC. NUM. COUNT: PATENT INFORMATION: PATENT NO. KIND DATE APPLICATION NO. DATE ----------_____ _____ -----WO 2000-US11085 20000425 <--WO 2000065064 A1 20001102 W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG US 6475492 B1 20021105 US 1999-300971 19990428 CA 2370493 AA 20001102 CA 2000-2370493 20000425 <--EP 2000-926350 20020116 EP 1171605 A1 20000425 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO 20031127 AU 2000-44893 20000425 AU 767955 B2 US 1999-300971 A2 19990428 PRIORITY APPLN. INFO.: WO 2000-US11085 W 20000425 ED Entered STN: 03 Nov 2000 A peptide consisting of an invariable 26-amino-acid-long region, named AB IR6, which is antigenically conserved among strains and species of the Borrelia burgdorferi sensu lato complex, and immunodominant in both human and nonhuman primate hosts, is described. This peptide is characterized by the sequence MKKDDQIAAAMVLRGMAKDGQFALKD. This peptide is useful for rapid and specific diagnosis of Lyme disease, as are proteins containing this peptide and nucleic acid sequences encoding this peptide and these proteins. Also provided is a novel ELISA, which is characterized by high sensitivity and specificity. ICM C12N015-31 IC C07K014-00; C07K014-20; C07K016-12; G01N033-68; A61K039-02; A61K038-00; C12N015-62 15-1 (Immunochemistry) CC Section cross-reference(s): 1 Genetic vectors IT (VlsE peptide- or peptides-containing fusion protein-encoding; vlsE-derived peptides and nucleic acids and vaccines for diagnosis and prevention of Lyme disease) ΤТ Cell (VlsE peptide- or peptides-containing fusion protein-producing; vlsE-derived peptides and nucleic acids and vaccines for diagnosis and prevention of Lyme disease) IT RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (for VlsE peptide or peptides-containing fusion protein; vlsE-derived peptides and nucleic acids and vaccines for diagnosis and prevention of Lyme disease) 303177-90-4 303177-89-1 303339-33-5 303339-34-6 303339-35-7

(unclaimed sequence; vlsE-derived peptides and nucleic acids and

RL: PRP (Properties)

TT

vaccines for diagnosis and prevention of Lyme disease)

REFERENCE COUNT: 7 THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L16 ANSWER 47 OF 125 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2000:772766 CAPLUS

DOCUMENT NUMBER: 133:330556

TITLE: Genome sequence and polypeptides of Pyrococcus abyssi

and their uses

INVENTOR(S): Forterre, Patrick; Thierry, Jean-Claude; Prieur,

> Daniel; Dietrich, Jacques; Lecompte, Odile; Querellou, Joel; Weissenbach, Jean; Saurin, William; Heilig, Roland; Flament, Didier; Raffin, Jean-Paul; Henneke,

Ghislaine; Gueguen, Yannick; Rolland, Jean-Luc

PATENT ASSIGNEE(S): Centre National de la Recherche Scientifique (CNRS),

Fr.; Institut Français de Recherche pour

l'Exploitation de la Mer - IFREMER

SOURCE: PCT Int. Appl., 1403 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent French LANGUAGE:

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PA	PATENT NO.				KIND DATE			APPLICATION NO.						DATE				
	O 2000065062			A2		2000 2002		WO 2000-FR1065						20000421 <				
	2000						2002											
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Entered STN: 03 Nov 2000 ED

The invention relates to the genome sequence of Pyrococcus abyssi strain AΒ Orsay, the 807 open reading frame nucleotide sequences coding for polypeptides of P. abyssi such as polypeptides involved in metabolism or in the replication process, in addition to vectors including said sequences and cells transformed by said vectors. Replication factor C (large and small forms resulting from intein splicing), PCNA (proliferating cell nuclear antigen), DNA polymerase II large and small subunits, replication factor A, and DNA polymerase I were isolated and characterized by recombinant cloning in Escherichia coli. The invention also relates to methods using said nucleic acids or polypeptides, especially biosynthesis methods or biodegrdn. methods for mols. of interest and to kits comprising said polypeptides.

ICM C12N015-31 IC

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C12N015-54; C12N015-55; C12N015-57; C12N015-60; C12N015-61;
          C12N015-62; C07K014-195; C07K019-00; C12N009-10; C12N009-12;
          C12N009-14; C12N009-16; C12N009-48; C12N009-88; C12N009-90;
          C12P001-00; C12P019-34; C12Q001-68; C12N001-21
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     3-3 (Biochemical Genetics)
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     Fusion proteins (chimeric proteins)
     RL: BPN (Biosynthetic preparation); BUU (Biological use, unclassified);
     BIOL (Biological study); PREP (Preparation); USES (Uses)
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        abyssi and their uses)
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    RL: BOC (Biological occurrence); BSU (Biological study, unclassified); BUU
     (Biological use, unclassified); PRP (Properties); BIOL (Biological study);
    OCCU (Occurrence); USES (Uses)
        (amino acid sequence; genome sequence and polypeptides of Pyrococcus
       abyssi and their uses)
L16 ANSWER 48 OF 125 CAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                      2000:772468 CAPLUS
                       133:331436
DOCUMENT NUMBER:
                       Nucleic acids and proteins of a rat ganglioside
TITLE:
                       GM1-specific \alpha1\rightarrow 2 fucosyltransferase and
                       synthetic uses
INVENTOR (S):
                       Holmes, Eric H.; Sherwood, Anne L.
PATENT ASSIGNEE(S):
                       Pacific Northwest Cancer Foundation, USA
SOURCE:
                       PCT Int. Appl., 91 pp.
                        CODEN: PIXXD2
DOCUMENT TYPE:
                       Patent
                        English
LANGUAGE:
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:
    PATENT NO.
                    KIND DATE
                                       APPLICATION NO.
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                                         ......
    WO 2000064464
                       A1 20001102 WO 1999-US7384
                                                              19990423 <--
        W: CA, JP
        RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL,
            PT, SE
PRIORITY APPLN. INFO.:
                                         WO 1999-US7384
                                                                19990423
    Entered STN: 03 Nov 2000
    A rat qanglioside GM1-specific \alpha 1 \rightarrow 2 fucosyltransferase (I) is
    disclosed. Nucleotide sequences of a rat I, amino acid sequences of its
    encoded protein (including peptide or polypeptide), and derivs. thereof
     are described. Also described are fragments (and derivs. and analogs
     thereof) which comprise a domain of rat I with catalytic activity.
    Methods of production of rat I and derivs. and analogs thereof (e.g. by
    recombinant means) are provided. Methods of inhibiting the function of
    rat I (e.g. by means of antisense RNA) are provided. Methods of com.
     scale use of the rat I in the production of fucosyl-saccharide compns. are
    described. Applications of these compns., e.g. as additives for human
    nutritive compns. or immunotherapeutics for cancer, are also disclosed.
    ICM A61K038-00
         C07H021-04; C12N015-00; C12N005-00; C12N006-02; C12P021-06;
     ICS
         A61K038-00; A01N037-18; C12Q001-68; A01N043-04; A61K039-00
    7-3 (Enzymes)
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Section cross-reference(s): 1, 3, 17

ED

AB

IC

CC

ITProteins, specific or class RL: BPN (Biosynthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses) (A, fusion protein with IqG binding domain of; nucleic acids and proteins of a rat ganglioside GM1-specific α 1 \rightarrow 2fucosyltransferase and its synthetic uses) IT RL: BOC (Biological occurrence); BSU (Biological study, unclassified); FFD (Food or feed use); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); OCCU (Occurrence); USES (Uses) (nucleotide sequence; nucleic acids and proteins of a rat ganglioside GM1-specific α1→2fucosyltransferase and its synthetic uses) REFERENCE COUNT: THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT L16 ANSWER 49 OF 125 CAPLUS COPYRIGHT 2006 ACS on STN ACCESSION NUMBER: 2000:742109 CAPLUS DOCUMENT NUMBER: 133:291985 TITLE: Cloning and cDNA and deduced amino acid sequences of 50 human secreted proteins INVENTOR(S): Ruben, Steven M.; Komatsoulis, George Human Genome Sciences, Inc., USA; Rosen, Craig A. PATENT ASSIGNEE(S): PCT Int. Appl., 505 pp. SOURCE: CODEN: PIXXD2 DOCUMENT TYPE: Patent LANGUAGE: English FAMILY ACC. NUM. COUNT: 1 PATENT INFORMATION: APPLICATION NO. KIND DATE PATENT NO. DATE --------------WO 2000061596 A1 20001019 WO 2000061596 C1 20020620 WO 2000-US8983 20000406 <--W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG CA 2371172 AA20001019 CA 2000-2371172 20000406 <--20001114 AU 2000-40723 20020227 EP 2000-920141 AU 2000040723 Α5 20000406 <--EP 1181303 A1 20000406 AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO JP 2003523726 T2 20030812 JP 2000-610865 20000406 PRIORITY APPLN. INFO.: US 1999-128703P P 19990409 US 2000-176068P P 20000120 WO 2000-US8983 W 20000406 Entered STN: 20 Oct 2000 ED AΒ The present invention relates to 50 novel human secreted proteins and isolated nucleic acids containing the coding regions of the genes encoding such proteins. Tissue distribution, sequence homologies, and preferred epitope sites are provided for the secreted proteins, as well as

chromosomal mapping of some of the genes. Also provided are vectors, host cells, antibodies, and recombinant methods for producing human secreted proteins in bacterial, insect, and mammalian cells. The invention further

relates to diagnostic and therapeutic methods useful for diagnosing and treating disorders related to these novel human secreted proteins. High-throughput screening assays are also provided for various putative activities of the secreted proteins.

IC ICM C07H021-02

> ICS C07H021-04; C12N001-20; C12N015-00; C12N015-09; C12N015-63; C12N015-70; C12N015-74

3-3 (Biochemical Genetics) CC

Section cross-reference(s): 6, 13, 63

IT Immunoglobulins

> RL: BPN (Biosynthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(fusion products; cloning and cDNA and deduced amino acid sequences of 50 human secreted proteins)

124025-38-3, Ubiquitin (human clone yep-HUBCEP52 precursor reduced) IT207354-71-0 212964-70-0, Protein (human clone BEC6 gene 191941-70-5 300625-85-8 246219-60-3 292158-94-2 300625-86-9 CREBL2) 300625-87-0 300625-88-1 300625-89-2 300625-90-5 300625-91-6 300625-92-7 300625-93-8 300625-94-9 300625-95-0 300625-98-3 300626-00-0 300625-96-1 300625-97-2 300625-99-4 300626-03-3 300682-14-8 300626-02-2 300626-04-4 300626-01-1 300693-57-6 300693-58-7 300693-59-8

300682-20-6 300682-18-2 300693-62-3 300693-60-1 300693-61-2 300693-63-4 300693-64-5 300693-65-6 300693-66-7 300693-67-8 300693-68-9

RL: PRP (Properties)

(unclaimed protein sequence; cloning and cDNA and deduced amino acid sequences of 50 human secreted proteins)

REFERENCE COUNT:

THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L16 ANSWER 50 OF 125 CAPLUS COPYRIGHT 2006 ACS on STN

2

ACCESSION NUMBER:

<2000:711033 CAPLUS

DOCUMENT NUMBER:

133:251261

TITLE:

-Fusion protein for immunoprophyaxis and

immunotherapy of venereal disease and cancer

INVENTOR(S): Zhou, Guoqing PATENT ASSIGNEE(S): Peop. Rep. China

Faming Zhuanli Shenging Gongkai Shuomingshu, 5 pp. SOURCE:

CODEN: CNXXEV

DOCUMENT TYPE: Patent LANGUAGE: Chinese

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
CN 1248631	Α	20000329	CN 1998-112264	19980924 <
PRIORITY APPLN. INFO.:			CN 1998-112264	19980924

Entered STN: 10 Oct 2000 ED

- The fusion protein is whole or part heat shock protein of Mycobacterium AB bovis var BCG connected with whole or part human papillary virus (HPV) antigen such as early-expressed proteins, and its N-terminal may be modified by several histidines. The fusion protein may be expressed in E. coli, yeast, or plant. The protein sequence of the recombinant fusion protein Hsp-E7 is presented. The fusion protein is used for immunol. prevention and treatment of fig wart, tumor and cancer induced by human papillary virus.
- IC ICM C12P021-00 ICS C07K019-00

CC 15-2 (Immunochemistry)

Section cross-reference(s): 1, 3, 16 STfusion protein human papillary virus; heat shock gene Mycobacterium fusion protein IT Transcription factors RL: BPN (Biosynthetic preparation); BIOL (Biological study); PREP (Preparation) (E7; fusion protein for immunoprophyaxis and immunotherapy of venereal disease and cancer) Heat-shock proteins ITRL: BPN (Biosynthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses) (HSP 65; fusion protein for immunoprophyaxis and immunotherapy of venereal disease and cancer) Proteins, specific or class TT RL: BPN (Biosynthetic preparation); BIOL (Biological study); PREP (Preparation) (early; fusion protein for immunoprophyaxis and immunotherapy of venereal disease and cancer) IT Escherichia coli Fermentation Human papillomavirus Human papillomavirus 16 Human papillomavirus 18 Immunotherapy Mycobacterium bovis Neoplasm Plant (Embryophyta) Protein sequences Sexually transmitted diseases Wart Yeast (fusion protein for immunoprophyaxis and immunotherapy of venereal disease and cancer) Fusion proteins (chimeric proteins) IT Heat-shock proteins RL: BPN (Biosynthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses) (fusion protein for immunoprophyaxis and immunotherapy of venereal disease and cancer) TT Peptides, biological studies RL: BUU (Biological use, unclassified); BIOL (Biological study); USES (Uses) (histidine tag; fusion protein for immunoprophyaxis and immunotherapy of venereal disease and cancer) 295371-00-5 IT RL: PRP (Properties) (amino acid sequence; fusion protein for immunoprophyaxis and immunotherapy of venereal disease and cancer) 71-00-1, Histidine, biological studies TT RL: BUU (Biological use, unclassified); BIOL (Biological study); USES (Uses) (tag; fusion protein for immunoprophyaxis and immunotherapy of venereal disease and cancer)